



US009192307B2

(12) **United States Patent**
Bates et al.

(10) **Patent No.:** **US 9,192,307 B2**
(45) **Date of Patent:** **Nov. 24, 2015**

(54) **SYSTEMS AND METHODS FOR
MINIMALLY-INVASIVE
OPTICAL-ACOUSTIC IMAGING**

(71) Applicant: **Vascular Imaging Corporation**,
Rancho Cordova, CA (US)

(72) Inventors: **Kenneth N. Bates**, Beaverton, OR (US);
Gil M. Vardi, Town & Country, MO
(US)

(73) Assignee: **Vascular Imaging Corporation**,
Rancho Cordova, CA (US)

(*) Notice: Subject to any disclaimer, the term of this
patent is extended or adjusted under 35
U.S.C. 154(b) by 0 days.

(21) Appl. No.: **14/280,327**

(22) Filed: **May 16, 2014**

(65) **Prior Publication Data**

US 2014/0254975 A1 Sep. 11, 2014

Related U.S. Application Data

(63) Continuation of application No. 13/779,985, filed on
Feb. 28, 2013, now Pat. No. 8,731,340, which is a
continuation of application No. 13/285,551, filed on
Oct. 31, 2011, now Pat. No. 8,391,652, which is a
(Continued)

(51) **Int. Cl.**
G02B 6/34 (2006.01)
A61B 1/00 (2006.01)
(Continued)

(52) **U.S. Cl.**
CPC **A61B 5/0095** (2013.01); **A61B 5/0097**
(2013.01); **A61B 5/02007** (2013.01);
(Continued)

(58) **Field of Classification Search**
CPC A61B 1/00117; A61B 1/00165–1/0017;
G02B 6/34
USPC 385/37; 600/478
See application file for complete search history.

(56) **References Cited**

U.S. PATENT DOCUMENTS

3,995,623 A 12/1976 Blake et al.
4,068,191 A 1/1978 Zemon et al.

(Continued)

FOREIGN PATENT DOCUMENTS

CA 2472877 A1 7/2003
DE 2363984 A1 6/1975

(Continued)

OTHER PUBLICATIONS

“U.S. Appl. No. 09/623,248, Corrected Notice of Allowance mailed
Jul. 11, 2003”, 8 pgs.

(Continued)

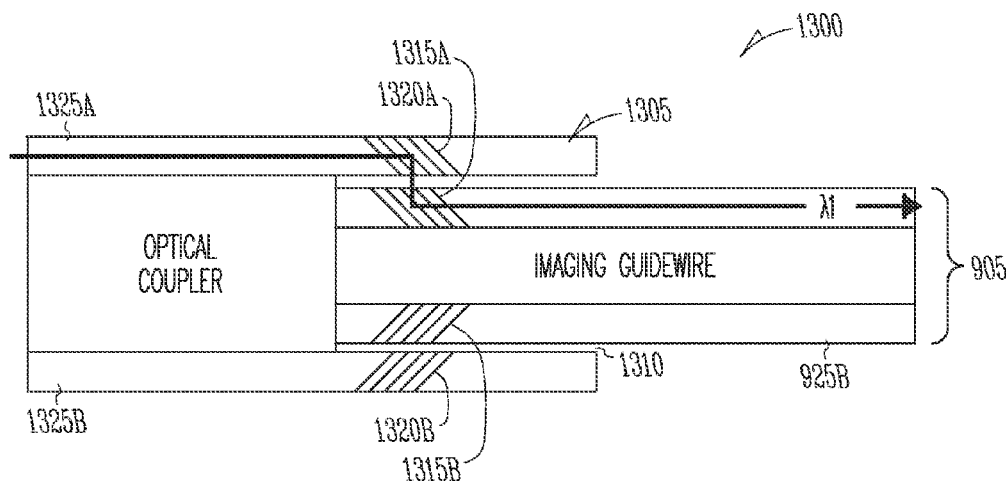
Primary Examiner — Jerry Rahlh

(74) *Attorney, Agent, or Firm* — Schwegman Lundberg &
Woessner, P.A.

(57) **ABSTRACT**

An imaging guidewire can include one or more optical fibers
communicating light along the guidewire. At or near its distal
end, one or more blazed or other fiber Bragg gratings (FBGs)
directs light to a photoacoustic transducer material that pro-
vides ultrasonic imaging energy. Returned ultrasound is
sensed by an FBG sensor. A responsive signal is optically
communicated to the proximal end of the guidewire, and
processed to develop a 2D or 3D image. In one example, the
guidewire outer diameter is small enough such that an intra-
vascular catheter can be passed over the guidewire. Tech-
niques for improving ultrasound reception include using a
high compliance material, resonating the ultrasound sensing
transducer, using an attenuation-reducing coating and/or
thickness, and/or using optical wavelength discrimination.
Techniques for improving the ultrasound generating trans-
ducer include using a blazed FBG, designing the photoacous-
tic material thickness to enhance optical absorption. Tech-
niques for distinguishing plaque or vulnerable plaque may be
used to enhance the displayed image.

8 Claims, 11 Drawing Sheets



Related U.S. Application Data

continuation of application No. 12/701,228, filed on Feb. 5, 2010, now Pat. No. 8,059,923, which is a continuation of application No. 12/263,978, filed on Nov. 3, 2008, now Pat. No. 7,660,492, which is a continuation of application No. 11/674,568, filed on Feb. 13, 2007, now Pat. No. 7,447,388, which is a continuation of application No. 10/266,082, filed on Oct. 7, 2002, now Pat. No. 7,245,789.

(51) Int. Cl.

A61B 5/00 (2006.01)
A61B 5/02 (2006.01)
A61B 8/12 (2006.01)
G01H 9/00 (2006.01)
G01N 29/24 (2006.01)
A61B 8/08 (2006.01)

(52) U.S. Cl.

CPC **A61B5/6876** (2013.01); **A61B 5/742** (2013.01); **A61B 8/12** (2013.01); **G01H 9/004** (2013.01); **G01N 29/24** (2013.01); **A61B 8/0833** (2013.01)

(56) References Cited

U.S. PATENT DOCUMENTS

4,076,379	A	2/1978	Chouinard	5,675,674	A	10/1997	Weis
4,115,753	A	9/1978	Shajenko	5,680,489	A	10/1997	Kersey
4,327,738	A	5/1982	Green et al.	5,682,897	A	11/1997	Pomeranz
4,473,065	A	9/1984	Bates	5,684,297	A	11/1997	Tardy et al.
4,522,193	A	6/1985	Bates	5,691,999	A	11/1997	Ball et al.
4,587,972	A	5/1986	Morantte, Jr.	5,693,043	A	12/1997	Kittrell et al.
4,887,605	A	12/1989	Angelsen et al.	5,700,236	A	12/1997	Sauer et al.
4,900,921	A	2/1990	Spillman, Jr.	5,704,361	A	1/1998	Seward et al.
4,917,097	A	4/1990	Proudian et al.	5,718,226	A	2/1998	Riza
4,946,238	A	8/1990	Sashin et al.	5,732,046	A	3/1998	O'Donnell et al.
5,007,705	A	4/1991	Morey et al.	5,748,564	A	5/1998	Pattanayak
5,070,882	A	12/1991	Bui et al.	5,774,610	A	6/1998	O'rourke et al.
5,095,911	A	3/1992	Pomeranz	5,779,643	A	7/1998	Lum et al.
5,099,090	A	3/1992	Allan et al.	5,779,644	A	7/1998	Eberle et al.
5,109,463	A	4/1992	Lee	5,830,145	A	11/1998	Tenhoff
5,135,295	A	8/1992	Jen et al.	5,852,233	A	12/1998	Arnold et al.
5,135,486	A	8/1992	Eberle et al.	5,857,974	A	1/1999	Eberle et al.
5,156,772	A	10/1992	Allan	5,865,178	A	2/1999	Yock
5,167,233	A	12/1992	Eberle et al.	5,872,879	A	2/1999	Hamm
5,183,048	A	2/1993	Eberle	5,873,835	A	2/1999	Hastings et al.
5,186,177	A	2/1993	O'Donnell et al.	5,876,344	A	3/1999	Baker et al.
5,226,847	A	7/1993	Thomas, III et al.	5,894,531	A	4/1999	Alcoz
5,240,004	A	8/1993	Walinsky et al.	5,921,931	A	7/1999	O'Donnell et al.
5,254,112	A	10/1993	Sinofsky et al.	5,938,609	A	8/1999	Pomeranz
5,290,275	A	3/1994	Kittrell et al.	5,938,615	A	8/1999	Eberle et al.
5,305,758	A	4/1994	Dietz et al.	5,944,687	A	8/1999	Benett et al.
5,325,860	A	7/1994	Seward et al.	5,953,477	A	9/1999	Wach et al.
5,353,262	A	10/1994	Yakymyshyn et al.	5,980,117	A	11/1999	Feuer et al.
5,368,037	A	11/1994	Eberle et al.	6,039,701	A	3/2000	Sliwa et al.
5,383,467	A	1/1995	Auer et al.	6,049,958	A	4/2000	Eberle et al.
5,400,788	A	3/1995	Dias et al.	6,057,927	A	5/2000	Levesque et al.
5,411,500	A	5/1995	Lafferty et al.	6,078,831	A	6/2000	Belef et al.
5,427,107	A	6/1995	Milo et al.	6,080,109	A	6/2000	Baker et al.
5,439,000	A	8/1995	Gunderson et al.	6,100,969	A	8/2000	Perez
5,453,575	A	9/1995	O'Donnell et al.	6,111,645	A	8/2000	Tearney et al.
5,469,520	A	11/1995	Morey et al.	6,123,673	A	9/2000	Eberle et al.
5,486,170	A	1/1996	Winston et al.	6,134,003	A	10/2000	Tearney et al.
5,493,113	A	2/1996	Dunphy et al.	6,210,339	B1	4/2001	Kiepen et al.
5,558,669	A	9/1996	Reynard	6,218,661	B1	4/2001	Schroeder et al.
5,558,699	A	9/1996	Nakashima et al.	6,222,970	B1	4/2001	Wach et al.
5,573,493	A	11/1996	Sauer et al.	6,228,078	B1	5/2001	Eggers et al.
5,582,171	A	12/1996	Chornenky et al.	6,238,347	B1	5/2001	Nix et al.
5,584,793	A	12/1996	Sauer et al.	6,248,076	B1	6/2001	White et al.
5,601,087	A	2/1997	Gunderson et al.	6,261,246	B1	7/2001	Pantages et al.
5,603,327	A	2/1997	Eberle et al.	6,282,011	B1	8/2001	Tearney et al.
5,615,675	A	4/1997	O'Donnell et al.	6,283,920	B1	9/2001	Eberle et al.
5,660,180	A	8/1997	Malinowski et al.	6,292,610	B1	9/2001	O'Rourke et al.
				6,306,096	B1	10/2001	Seward et al.
				6,315,732	B1	11/2001	Suorsa et al.
				6,330,383	B1	12/2001	Cai et al.
				6,379,325	B1	4/2002	Benett et al.
				6,416,234	B1	7/2002	Wach et al.
				6,421,164	B2	7/2002	Tearney et al.
				6,494,836	B2	12/2002	Ogawa
				6,538,807	B2	3/2003	Kakui et al.
				6,575,965	B1	6/2003	Fitch et al.
				6,585,660	B2	7/2003	Dorando et al.
				6,611,633	B1	8/2003	Vohra et al.
				6,618,916	B1	9/2003	Eberle et al.
				6,659,957	B1	12/2003	Vardi et al.
				6,779,257	B2	8/2004	Kiepen et al.
				6,783,494	B2	8/2004	Ogawa
				6,819,845	B2	11/2004	Lee et al.
				6,839,496	B1	1/2005	Mills et al.
				6,907,163	B2	6/2005	Lewis
				6,938,474	B2	9/2005	Melväs
				6,948,859	B2	9/2005	Anderson
				6,984,819	B2	1/2006	Ogawa
				7,082,238	B2	7/2006	Nishimura
				7,097,620	B2	8/2006	Corl et al.
				7,184,148	B2	2/2007	Alphonse
				7,190,464	B2	3/2007	Alphonse
				7,242,480	B2	7/2007	Alphonse
				7,242,832	B2	7/2007	Carlin et al.
				7,245,789	B2	7/2007	Bates et al.
				7,417,740	B2	8/2008	Alphonse et al.
				7,447,388	B2	11/2008	Bates et al.
				7,527,594	B2	5/2009	Vardi et al.
				7,599,588	B2	10/2009	Eberle et al.

(56)

References Cited**U.S. PATENT DOCUMENTS**

7,660,492	B2	2/2010	Bates et al.	
7,753,852	B2	7/2010	Maschke	
7,881,573	B2	2/2011	Eberle et al.	
8,059,923	B2	11/2011	Bates et al.	
8,218,927	B2	7/2012	Chang et al.	
8,320,723	B2	11/2012	Eberle et al.	
8,391,652	B2	3/2013	Bates et al.	
8,560,048	B2	10/2013	Eberle et al.	
8,731,340	B2	5/2014	Bates et al.	
8,861,908	B2	10/2014	Eberle et al.	
8,926,519	B2	1/2015	Vardi et al.	
8,968,376	B2	3/2015	Wells et al.	
2001/0021843	A1	9/2001	Bosselmann et al.	
2001/0043772	A1	11/2001	Sorin	
2001/0046352	A1*	11/2001	Ohta et al.	385/50
2002/0039463	A1	4/2002	Degertekin et al.	
2002/0041735	A1	4/2002	Cai et al.	
2002/0059827	A1	5/2002	Smith	
2002/0166955	A1	11/2002	Ogawa	
2003/0026546	A1	2/2003	Deliwala	
2003/0053774	A1	3/2003	Blomquist et al.	
2003/0060707	A1	3/2003	Ogawa	
2003/0118297	A1	6/2003	Dunphy et al.	
2003/0181802	A1	9/2003	Ogawa	
2003/0187369	A1	10/2003	Lewis et al.	
2004/0067000	A1	4/2004	Bates et al.	
2004/0116800	A1	6/2004	Helfer et al.	
2004/0182315	A1	9/2004	Laflamme, Jr. et al.	
2005/0020926	A1	1/2005	Wiklof et al.	
2005/0121734	A1	6/2005	Degertekin et al.	
2005/0131289	A1	6/2005	Aharoni et al.	
2005/0238292	A1	10/2005	Barnes et al.	
2006/0067616	A1	3/2006	Kanji et al.	
2007/0116408	A1	5/2007	Eberle et al.	
2007/0123776	A1	5/2007	Aharoni et al.	
2007/0133925	A1	6/2007	Bates et al.	
2007/0291275	A1	12/2007	Diamond	
2008/0077225	A1	3/2008	Carlin et al.	
2008/0114254	A1	5/2008	Matcovitch et al.	
2008/0119739	A1	5/2008	Vardi et al.	
2009/0059727	A1	3/2009	Bates et al.	
2010/0014810	A1	1/2010	Eberle et al.	
2010/0087732	A1	4/2010	Eberle et al.	
2010/0135111	A1	6/2010	Bates et al.	
2011/0123154	A1	5/2011	Eberle et al.	
2012/0108943	A1	5/2012	Bates et al.	
2013/0148933	A1	6/2013	Eberle et al.	
2013/0178729	A1	7/2013	Bates et al.	
2014/0142414	A1	5/2014	Eberle et al.	
2014/0180031	A1	6/2014	Anderson	
2014/0180034	A1	6/2014	Hoseit et al.	
2014/0200438	A1	7/2014	Millett et al.	
2015/0045645	A1	2/2015	Eberle et al.	

FOREIGN PATENT DOCUMENTS

EP	0478410	A1	4/1992
EP	1059878	A1	12/2000
EP	1152240	A2	11/2001
GB	2270159	A	3/1994
JP	59158699	A	9/1984
JP	6354151	A	3/1988
JP	63102421	A	5/1988
JP	2503279	A	10/1990
JP	04355415	A	12/1992
JP	05015536	A2	1/1993
JP	05034550	A	2/1993
JP	5220152	A	8/1993
JP	06003550	A	1/1994
JP	08112289	A2	5/1996
JP	9010215	A	1/1997
JP	09187513	A	7/1997
JP	10073742	A	3/1998
JP	10505920	A	6/1998

JP	10507036	A	7/1998
JP	10510364	A	10/1998
JP	11194280	A	7/1999
JP	11243596	A	9/1999
JP	11514432	A	12/1999
JP	2000508939	A	7/2000
JP	2001091785	A	4/2001
JP	2002514455	A	5/2002
JP	2003232964	A	8/2003
JP	2004085756	A	3/2004
JP	2004177549	A	6/2004
JP	2005079177	A	3/2005
JP	4733982	B2	4/2011
WO	WO-8809150	A1	1/1988
WO	WO-8907419	A1	8/1989
WO	WO-9739691	A1	10/1997
WO	WO-9958059	A1	11/1999
WO	WO-0049938	A1	8/2000
WO	WO-0121244	A1	3/2001
WO	WO-0219898	A3	3/2002
WO	WO-02054948	A1	7/2002
WO	WO-02075404	A1	9/2002
WO	WO-03057061	A1	7/2003
WO	WO-2004008070	A2	1/2004
WO	WO-2004029667	A2	4/2004
WO	WO-2004032746	A2	4/2004
WO	WO-2004077100	A2	9/2004
WO	WO-2004090484	A2	10/2004
WO	WO-2007062050	A2	5/2007
WO	WO-2007062050	A3	5/2007
WO	WO-2010039950	A1	4/2010

OTHER PUBLICATIONS

“U.S. Appl. No. 09/623,248, Examiner Interview Summary filed Nov. 6, 2003”, 1 pg.

“U.S. Appl. No. 09/623,248, Notice of Allowance mailed Jan. 13, 2003”, 7 pgs.

“U.S. Appl. No. 09/623,248, Notice of Allowance mailed Jun. 2, 2003”, 7 pgs.

“U.S. Appl. No. 09/623,248, Request for Continued Examination filed Apr. 14, 2003”, 6 pgs.

“U.S. Appl. No. 09/623,248, Supplemental Notice of Allowability mailed Oct. 7, 2003”, 6 pgs.

“U.S. Appl. No. 10/266,082, Non Final Office Action mailed Oct. 4, 2005”, 5 pgs.

“U.S. Appl. No. 10/266,082, Non-Final Office Action mailed Apr. 5, 2006”, 7 pgs.

“U.S. Appl. No. 10/266,082, Notice of Allowance mailed Jan. 12, 2005”, 5 pgs.

“U.S. Appl. No. 10/266,082, Notice of Allowance mailed Jan. 16, 2007”, 5 pgs.

“U.S. Appl. No. 10/266,082, Notice of Allowance mailed Mar. 22, 2007”, 5 pgs.

“U.S. Appl. No. 10/266,082, Notice of Allowance mailed Aug. 26, 2004”, 8 pgs.

“U.S. Appl. No. 10/266,082, Notice of Allowance mailed Sep. 22, 2006”, 4 pgs.

“U.S. Appl. No. 10/266,082, Response filed Jan. 3, 2006 to Non Final Office Action mailed Oct. 4, 2005”, 10 pgs.

“U.S. Appl. No. 10/266,082, Response filed Jul. 5, 2006 to Non Final Office Action mailed Apr. 5, 2006”, 9 pgs.

“U.S. Appl. No. 10/266,082, Response filed Jul. 28, 2004 to Restriction Requirement Jun. 29, 2004”, 2 pgs.

“U.S. Appl. No. 10/266,082, Restriction Requirement mailed Jun. 29, 2004”, 5 pgs.

“U.S. Appl. No. 10/685,226, Advisory Action mailed Apr. 17, 2006”, 3 pgs.

“U.S. Appl. No. 10/685,226, Final Office Action mailed Jan. 13, 2006”, 6 pgs.

“U.S. Appl. No. 10/685,226, Non Final Office Action mailed Jan. 18, 2007”, 6 pgs.

“U.S. Appl. No. 10/685,226, Non Final Office Action mailed Jun. 15, 2005”, 4 pgs.

(56)

References Cited**OTHER PUBLICATIONS**

"U.S. Appl. No. 10/685,226, Non Final Office Action mailed Jul. 24, 2006", 7 pgs.
 "U.S. Appl. No. 10/685,226, Notice of Allowance mailed Oct. 18, 2007", 5 pgs.
 "U.S. Appl. No. 10/685,226, Preliminary Amendment filed OPct. 14, 2003", 1 pg.
 "U.S. Appl. No. 10/685,226, Response filed Mar. 13, 2006 to Final Office Action mailed Jan. 13, 2006", 12 pgs.
 "U.S. Appl. No. 10/685,226, Response filed Apr. 18, 2007 to Non-Final Office Action mailed Jan. 18, 2007", 10 pgs.
 "U.S. Appl. No. 10/685,226, Response filed Oct. 14, 2005 to Non Final Office Action mailed Jun. 15, 2005", 15 pgs.
 "U.S. Appl. No. 10/685,226, Response filed Oct. 23, 2006 to Non Final Office Action mailed Jul. 24, 2006", 9 pgs.
 "U.S. Appl. No. 11/285,499, Advisory Action mailed Jun. 24, 2008", 4 pgs.
 "U.S. Appl. No. 11/285,499, Decision on Pre-Appeal Brief Request mailed Sep. 9, 2008", 2 pgs.
 "U.S. Appl. No. 11/285,499, Examiner Interview Summary mailed May 18, 2009", 4 pgs.
 "U.S. Appl. No. 11/285,499, Final Office Action mailed Jan. 25, 2008", 7 pgs.
 "U.S. Appl. No. 11/285,499, Non Final Office Action mailed May 16, 2007", 13 pgs.
 "U.S. Appl. No. 11/285,499, Non-Final Office Action mailed Nov. 13, 2008", 7 pgs.
 "U.S. Appl. No. 11/285,499, Notice of Allowance mailed May 27, 2009", 6 pgs.
 "U.S. Appl. No. 11/285,499, Pre-Appeal Brief Request filed Jul. 24, 2008", 5 pgs.
 "U.S. Appl. No. 11/285,499, Response filed Feb. 15, 2007 to Restriction Requirement mailed Jan. 26, 2007", 12 pgs.
 "U.S. Appl. No. 11/285,499, Response filed Apr. 13, 2009 to Non Final Office Action mailed Nov. 13, 2008", 14 pgs.
 "U.S. Appl. No. 11/285,499, Response filed May 27, 2008 to Final Office Action mailed Jan. 25, 2008", 9 pgs.
 "U.S. Appl. No. 11/285,499, Response filed Oct. 16, 2007 to Non-Final Office Action mailed May 16, 2007", 11 pgs.
 "U.S. Appl. No. 11/285,499, Restriction Requirement mailed Jan. 26, 2007", 4 pgs.
 "U.S. Appl. No. 11/674,568, Non-Final Office Action mailed Jan. 7, 2008", 6 pgs.
 "U.S. Appl. No. 11/674,568, Notice of Allowance mailed Jun. 25, 2008", 4 pgs.
 "U.S. Appl. No. 11/674,568, Response filed Apr. 21, 2008 to Non Final Office Action mailed Jan. 7, 2008", 7 pgs.
 "U.S. Appl. No. 11/674,568, Response filed Oct. 16, 2007 to Restriction Requirement mailed Sep. 17, 2007", 7 pgs.
 "U.S. Appl. No. 11/674,568, Restriction Requirement mailed Sep. 17, 2007", 6 pgs.
 "U.S. Appl. No. 12/020,736, Advisory Action mailed Feb. 13, 2014", 3 pgs.
 "U.S. Appl. No. 12/020,736, Examiner Interview Summary mailed May 8, 2014", 3 pgs.
 "U.S. Appl. No. 12/020,736, Final Office Action mailed Oct. 12, 2012", 15 pgs.
 "U.S. Appl. No. 12/020,736, Final Office Action mailed Oct. 25, 2013", 17 pgs.
 "U.S. Appl. No. 12/020,736, Non Final Office Action mailed Jun. 4, 2013", 17 pgs.
 "U.S. Appl. No. 12/020,736, Non Final Office Action mailed Sep. 30, 2011", 13 pgs.
 "U.S. Appl. No. 12/020,736, Non Final Office Action mailed Dec. 10, 2010", 10 pgs.
 "U.S. Appl. No. 12/020,736, Response filed Mar. 13, 2014 to Advisory Action mailed Feb. 13, 2014", 14 pgs.
 "U.S. Appl. No. 12/020,736, Response filed Mar. 30, 2012 to Non Final Office Action mailed Sep. 30, 2011", 17 pgs.

"U.S. Appl. No. 12/020,736, Response filed Apr. 12, 2013 to Final Office Action mailed Oct. 12, 2012", 17 pgs.
 "U.S. Appl. No. 12/020,736, Response filed Jun. 10, 2011 to Non-Final Office Action mailed Dec. 10, 2010", 9 pgs.
 "U.S. Appl. No. 12/020,736, Response filed Oct. 4, 2013 to Non Final Office Action mailed Jun. 4, 2013", 18 pgs.
 "U.S. Appl. No. 12/020,736, Response filed Dec. 18, 2013 to Final Office Action mailed Oct. 25, 2013", 11 pgs.
 "U.S. Appl. No. 12/020,736, Supplemental Response filed May 16, 2014 to Final Office Action mailed Oct. 25, 2013", 13 pgs.
 "U.S. Appl. No. 12/263,978, Notice of Allowance mailed Sep. 22, 2009", 6 pgs.
 "U.S. Appl. No. 12/571,724, Examiner Interview Summary mailed May 22, 2013", 3 pgs.
 "U.S. Appl. No. 12/571,724, Final Office Action mailed Jan. 4, 2013", 13 pgs.
 "U.S. Appl. No. 12/571,724, Non Final Office Action mailed Apr. 18, 2012", 11 pgs.
 "U.S. Appl. No. 12/571,724, Notice of Allowance mailed Jun. 11, 2013", 6 pgs.
 "U.S. Appl. No. 12/571,724, Response filed Jun. 4, 2013 to Final Office Action mailed Jan. 4, 2013", 13 pgs.
 "U.S. Appl. No. 12/571,724, Response filed Oct. 16, 2012 to Non Final Office Action mailed Apr. 18, 2012", 12 pgs.
 "U.S. Appl. No. 12/572,511, Non-Final Office Action mailed Jun. 1, 2010", 8 pgs.
 "U.S. Appl. No. 12/572,511, Notice of Allowance mailed Sep. 23, 2010", 6 pgs.
 "U.S. Appl. No. 12/572,511, Response filed May 17, 2010 to Restriction Requirement mailed May 10, 2010", 7 pgs.
 "U.S. Appl. No. 12/572,511, Response filed Sep. 1, 2010 to Non Final Office Action mailed Jun. 1, 2010", 9 pgs.
 "U.S. Appl. No. 12/572,511, Restriction Requirement mailed May 10, 2010", 5 pgs.
 "U.S. Appl. No. 12/701,228, Notice of Allowance mailed Apr. 22, 2011", 8 pgs.
 "U.S. Appl. No. 12/701,228, Notice of Allowance mailed Jun. 27, 2011", 5 pgs.
 "U.S. Appl. No. 13/017,354, Non Final Office Action mailed Jan. 9, 2012", 8 pgs.
 "U.S. Appl. No. 13/017,354, Non Final Office Action mailed Jun. 24, 2011", 8 pgs.
 "U.S. Appl. No. 13/017,354, Notice of Allowance mailed Jul. 24, 2012", 7 pgs.
 "U.S. Appl. No. 13/017,354, Response filed Jul. 5, 2012 to Non Final Office Action mailed Jan. 9, 2012", 15 pgs.
 "U.S. Appl. No. 13/017,354, Response filed Sep. 26, 2011 to Non-Final Office Action mailed Jun. 24, 2011", 16 pgs.
 "U.S. Appl. No. 13/285,551, Non Final Office Action mailed Apr. 12, 2012", 5 pgs.
 "U.S. Appl. No. 13/285,551, Notice of Allowance mailed Nov. 5, 2012", 7 pgs.
 "U.S. Appl. No. 13/285,551, Response filed Oct. 12, 2012 to Non Final Office Action mailed Apr. 12, 2012", 8 pgs.
 "U.S. Appl. No. 13/685,048, Non Final Office Action mailed Aug. 29, 2013", 10 pgs.
 "U.S. Appl. No. 13/685,048, Notice of Allowance mailed Feb. 20, 2014", 9 pgs.
 "U.S. Appl. No. 13/685,048, Notice of Allowance mailed Jun. 9, 2014", 9 pgs.
 "U.S. Appl. No. 13/685,048, Notice of Allowance mailed Nov. 22, 2013", 10 pgs.
 "U.S. Appl. No. 13/685,048, Response filed Nov. 6, 2013 to Non Final Office Action mailed Aug. 29, 2013", 11 pgs.
 "U.S. Appl. No. 13/779,985, Non Final Office Action mailed Apr. 25, 2013", 7 pgs.
 "U.S. Appl. No. 13/779,985, Notice of Allowance mailed Jan. 9, 2014", 16 pgs.
 "U.S. Appl. No. 13/779,985, Response filed Sep. 25, 2013 to Non Final Office Action mailed Apr. 25, 2013", 12 pgs.
 "U.S. Appl. No. 14/053,421, Non Final Office Action mailed Jun. 20, 2014", 13 pgs.

(56)

References Cited**OTHER PUBLICATIONS**

"Canadian Application Serial No. 2,348,580, Office Action mailed Feb. 20, 2007", 2 pgs.

"Canadian Application Serial No. 2,348,580, Response filed Aug. 16, 2007 to Office Action mailed Feb. 20, 2007", 7 pgs.

"Canadian Application Serial No. 2,501,048, Office Action mailed Aug. 26, 2013", 2 pgs.

"Canadian Application Serial No. 2,501,048, Office Action mailed Nov. 29, 2011", 3 pgs.

"Canadian Application Serial No. 2,501,048, Office Action mailed Jun. 4, 2012", 3 pgs.

"Canadian Application Serial No. 2,501,048, Response filed Feb. 11, 2014 to Office Action mailed Aug. 26, 2013", 10 pgs.

"Canadian Application Serial No. 2,501,048, Response filed Dec. 4, 2012 to Office Action mailed Jun. 4, 2012", 17 pgs.

"Canadian Application Serial No. 2,630,662, Office Action mailed Sep. 6, 2013", 3 pgs.

"Canadian Application Serial No. 2,630,662, Response filed Mar. 6, 2014 to Office Action mailed Sep. 6, 2013", 15 pgs.

"Canadian Application Serial No. 2,501,048, Response filed May 8, 2012 to Office Action mailed Nov. 29, 2011", 11 pgs.

"European Application Serial No. 03756904.3 Office Action mailed Nov. 3, 2009", 2 pgs.

"European Application Serial No. 03756904.3, Office Action mailed May 31, 2013", 2 pgs.

"European Application Serial No. 03756904.3, Office Action mailed Aug. 1, 2011", 4 pgs.

"European Application Serial No. 03756904.3, Response filed Oct. 10, 2013 to Office Action mailed May 31, 2013", 17 pgs.

"European Application Serial No. 03756904.3, Response filed Feb. 13, 2012 to Office Action mailed Aug. 1, 2011", 17 pgs.

"European Application Serial No. 03756904.3, Response filed May 13, 2010 to Office Action mailed Nov. 3, 2009", 23 pgs.

"European Application Serial No. 05024287.4, Office Action mailed Feb. 9, 2012", 4 pgs.

"European Application Serial No. 05024287.4, Office Action mailed Sep. 22, 2006", 1 pg.

"European Application Serial No. 05024287.4, Office Action mailed Sep. 25, 2012", 1 pg.

"European Application Serial No. 05024287.4, Response filed Nov. 23, 2012 to Office Action mailed Sep. 25, 2012", 9 pgs.

"European Application Serial No. 05024287.4, Response filed Jun. 6, 2007 to Office Action mailed Sep. 22, 2006", 8 pgs.

"European Application Serial No. 05024287.4, Search Report mailed Jan. 3, 2006", 7 pgs.

"European Application Serial No. 06838195.3, Office Action mailed Jul. 27, 2009", 4 pgs.

"European Application Serial No. 06838195.3, Response filed Feb. 4, 2010 to Office Action mailed Jul. 27, 2009", 8 pgs.

"European Application Serial No. 06838195.3, Office Action mailed Dec. 7, 2012", 7 pgs.

"European Application Serial No. 06838195.3, Response filed Jan. 15, 2013 to Office Action mailed Dec. 7, 2012", 1 pg.

"European Application Serial No. 09793238.8, Amendment Filed Apr. 29, 2011", 19 pgs.

"European Application Serial No. 99950325.3, Amendment filed Feb. 7, 2005", 11 pgs.

"European Application Serial No. 99950325.3, European Search Report mailed Mar. 8, 2004", 3 pgs.

"European Application Serial No. 99950325.3, Office Action mailed Jul. 28, 2004", 3 pgs.

"International Application Serial No. PCT/US03/31280, Demand and Response filed May 6, 2004 to Partial Search Report mailed Dec. 2, 2004", 13 pgs.

"International Application Serial No. PCT/US03/31280, International Preliminary Examination Report mailed Feb. 1, 2005", 16 pgs.

"International Application Serial No. PCT/US03/31280, International Search Report mailed Jul. 19, 2004", 5 pgs.

"International Application Serial No. PCT/US03/31280, Invitation to Pay Additional Fees and Partial Search Report mailed Feb. 12, 2004", 10 pgs.

"International Application Serial No. PCT/US2003/031280, Response filed Jan. 19, 2005 to Written Opinion mailed Jan. 14, 2005", 14 pgs.

"International Application Serial No. PCT/US2003/031280, Written Opinion mailed Jan. 14, 2005", 7 pgs.

"International Application Serial No. PCT/US2006/045080, International Preliminary Report on Patentability mailed Jun. 5, 2008", 10 pgs.

"International Application Serial No. PCT/US2006/045080, International Search Report and Written Opinion mailed May 16, 2007", 16 pgs.

"International Application Serial No. PCT/US2006/045080, Partial International Search Report mailed Mar. 9, 2007", 3 pgs.

"International Application Serial No. PCT/US2009/059218, International Preliminary Report on Patentability mailed Apr. 5, 2011", 9 pgs.

"International Application Serial No. PCT/US2009/059218, International Search Report mailed Feb. 12, 2010", 3 pgs.

"International Application Serial No. PCT/US2009/059218, Written Opinion mailed Apr. 2, 2011", 8 pgs.

"International Application Serial No. PCT/US99/04913, International Search Report mailed May 28, 1999", 1 pg.

"Japanese Application Serial No. 2000-547913, Office Action mailed Feb. 24, 2009", w/ English translation, 7 pgs.

"Japanese Application Serial No. 2000-547913, Office Action mailed Jun. 23, 2009", w/ English translation, 8 pgs.

"Japanese Application Serial No. 2000-547913, Response filed May 22, 2009 to Office Action mailed Feb. 24, 2009", w/English claims, 9 pgs.

"Japanese Application Serial No. 2000-547913, Response filed Dec. 18, 2009 to Office Action mailed Jun. 23, 2009", w/ English translation of claims, 12 pgs.

"Japanese Application Serial No. 2004-543092, Response filed Nov. 15, 2010 to Office Action mailed Jul. 13, 2010", w/English claims, 24 pgs.

"Japanese Application Serial No. 2004-543092, Office Action mailed Jul. 13, 2010", w/English translation, 3 pgs.

"Japanese Application Serial No. 2004-543092, Office Action mailed Nov. 17, 2009", w/English translation, 8 pgs.

"Japanese Application Serial No. 2004-543092, Response filed May 17, 2010 to Office Action mailed Nov. 17, 2009", 18 pgs.

"Japanese Application Serial No. 2008-541424, Amendment filed Nov. 16, 2009", w/English claims, 10 pgs.

"Japanese Application Serial No. 2008-541424, Office Action mailed Sep. 3, 2013", w/English translation, 6 pgs.

"Japanese Application Serial No. 2008-541424, Office Action mailed Oct. 30, 2012", w/ English translation, 8 pgs.

"Japanese Application Serial No. 2008-541424, Response filed Jan. 29, 2013 to Office Action mailed Oct. 30, 2012", w/English claims, 15 pgs.

"Japanese Application Serial No. 2008-541424, Response filed Oct. 29, 2013 to Office Action mailed Sep. 13, 2013", w/English claims, 9 pgs.

"Japanese Application Serial No. 2010-113577, Amendment filed Nov. 15, 2010", w/English claims, 16 pgs.

"Japanese Application Serial No. 2010-113577, Office Action mailed Feb. 14, 2012", w/English translation, 4 pgs.

"Japanese Application Serial No. 2010-113577, Office Action mailed Aug. 14, 2012", w/English translation, 6 pgs.

"Japanese Application Serial No. 2010-113577, Response filed Feb. 14, 2013 to Office Action mailed Aug. 14, 2012", 8 pgs.

"Japanese Application Serial No. 2010-113577, Response filed May 2, 2012 to Office Action mailed Feb. 14, 2012", w/English claims, 9 pgs.

"Japanese Application Serial No. 2004-543092, Notice of Allowance mailed Apr. 5, 2011", w/English translation, 2 pgs.

"Optical Review", vol. 4 No. 6, (1997), 1 pg.

"Tissue Characterization through Ultrasonic Backscatter", [Online]. Retrieved from the Internet: <URL: <http://www.brl.uiuc.edu/Projects/backscatter.htm>>, (retrieved Sep. 25, 2002), 5 pgs.

(56)

References Cited**OTHER PUBLICATIONS**

- Bates, K. N., "A High Acuity 3D Acoustic Imaging System", Proceedings, 1995 IEEE Ultrasonics Symposium, 2, (Nov. 7-10, 1995), 1245-1250.
- Bates, K. N., "A One Dimensional Phased Array Imaging System", Ph.D. Dissertation, Applied Physics, Stanford University, (1982), 186 pgs.
- Bates, K. N., et al., "PEOATS and ESOATS", IEEE Ultrasonics Symposium Proceedings, 1979, (Sep. 26-28, 1979), 189-194.
- Bates, K. N., "Tolerance Analysis for Phased Arrays", Acoustic Imaging, 9, (1980), 239-262.
- Bates, Kenneth N., "A high acuity 3-D acoustic imaging system", Proceedings, 1995 IEEE Ultrasonics Symposium, 2, (Nov. 7-10, 1995), 1245-1250.
- Bates, Kenneth N., et al., "Digitally Controlled Electronically Scanned and Focused Ultrasonic Imaging System", IEEE Ultrasonics Symposium Proceedings, 1979, (Sep. 26-28, 1979), 216-220.
- Blotekjaer, K., "Theoretical concepts of a novel Vernier-based fringe-counting fibre optic sensor", IEE Proceedings, Optoelectronics, 144(3), (Jun. 1997), 126-129.
- Brady, G. P., et al., "Simultaneous measurement of strain and temperature using the first-and second-order diffraction wavelengths of Bragg gratings", IEE Proceedings, Optoelectronics, 144(3), (Jun. 1997), 156-161.
- Buma, T., et al., "A high frequency ultrasound array element using thermoelastic expansion in PDMS", Proceedings of the 2001 IEEE Ultrasonics Symposium, 2, (Oct. 7-10, 2001), 1143-1146.
- Buma, T., et al., "A high-frequency, 2-D array element using thermoelastic expansion in PDMS", IEEE Transactions on Ultrasonics, Ferroelectrics and Frequency Control, 50(9), (Sep. 2003), 1161-1176.
- Buma, T., et al., "High Frequency Ultrasonic Imaging Using Optoacoustic Arrays", Proceedings of the Proceeding of the 2002 IEEE Ultrasonics Symposium, 1, Invited paper, (Oct. 8-11, 2002), 571-580.
- Buma, T., et al., "High Frequency Ultrasound Array Element using Thermoelastic Expansion in an Elastomeric Film", Applied Physics Letters, 79(4), (Jul. 23, 2001), 548-550.
- Buma, T., et al., "High-frequency ultrasound imaging using optoacoustic arrays", Proceedings of the SPIE—The International Society for Optical Engineering, 4687, (2002), 99-109.
- Buma, T., et al., "Thermoelastic Expansion versus Piezoelectricity for High Frequency 2-D Arrays", IEEE Transactions on Ultrasonics, Ferroelectrics and Frequency Control, 50(8), (Aug. 2003), 1065-1068.
- Buma, T., et al., "Thermoelastic Generation of Continuous Lamb Waves for Microfluidic Devices", Proceeding of the 2003 IEEE Ultrasonics Symposium, (2003), 150-153.
- Buma, T., et al., "Thermoelastic Generation of Ultrasound Using an Erbium Doped Fiber Amplifier", Proceeding of the 1999 IEEE Ultrasonics Symposium, 2, (Oct. 17-20, 1999), 1253-1256.
- Davis, M. A., et al., "Simultaneous measurement of temperature and strain using fibre Bragg gratings and Brillouin scattering", IEE Proceedings, Optoelectronics, 144(3), (Jun. 1997), 151-155.
- Feed, R., et al., "Advances in high resolution distributed temperature sensing using the time-correlated single photon counting technique", IEE Proceedings, Optoelectronics, 144(3), (Jun. 1997), 183-188.
- Furstenau, N., et al., "Extrinsic Fabry-Perot interferometer vibration and acoustic sensor systems for airport ground traffic monitoring", IEE Proceedings, Optoelectronics, 144(3), (Jun. 1997), 134-144.
- Hamilton, J. D., et al., "An active optical detector for high frequency ultrasound imaging", Proceedings of the 1997 IEEE Ultrasonics Symposium, 1, (Oct. 5-8, 1997), 753-756.
- Hamilton, J. D., et al., "High frequency optoacoustic arrays using etalon detection", IEEE Transactions on Ultrasonics, Ferroelectrics and Frequency Control, 47(1), (Jan. 2000), 160-169.
- Hamilton, J. D., et al., "High Frequency Ultrasound Imaging Using an Active optical Detector", IEEE Transactions on Ultrasonics, Ferroelectrics and Frequency Control UFFC-45, (1998), 719-727.
- Hamilton, J. D., et al., "High Frequency Ultrasound Imaging with Optical Arrays", IEEE Transactions on Ultrasonics, Ferroelectrics and Frequency Control, 45(1), (Jan. 1998), 216-235.
- Hamilton, J. D., et al., "Optical Arrays for High Frequency Ultrasound Imaging", Proceedings of the 1996 IEEE Ultrasonics Symposium, (1996), 1419-1422.
- Karl, W. Clem, "Multi-Sensor Fusion for Atherosclerotic Plaque Characterization", Boston University—MDSP, [Online] Retrieved from the internet: <<http://www.censsis.neu/Education/StudentResearch/2001/posters/weisenselljrl.pdf>>, (Retrieved Sep. 25, 2002), 15 pgs.
- Komiyama, N., et al., "Tissue Characterization of Atherosclerotic Plaques by Intravascular Ultrasound Radiofrequency Signal Analysis: An In Vitro Study of Human Coronary Arteries", American Heart Journal, 140(4), (Oct. 2000), 565-574.
- Krass, S., et al., "P3.4 Pattern Recognition Algorithms for Tissue Characterization in Intracoronary Ultrasound Imaging: Test Data Set and Results of Computerized Texture Analysis", 2nd Medical Clinic, Univ-Mainz, Germany, [online]. Retrieved from the Internet: <URL: http://www.uni-mainz.de/Cardio/incis/Data/p3_4.htm>, (Accessed on Sep. 25, 2002), 3 pgs.
- Lockey, R. A., et al., "Multicomponent time-division-multiplexed optical fibre laser Doppler anemometry", IEE Proceedings, Optoelectronics, 144(3), (Jun. 1997), 168-175.
- MacPherson, W. N., et al., "Phase demodulation in optical fibre Fabry-Perot sensors with inexact phase steps", IEE Proceedings, Optoelectronics, 144(3), (Jun. 1997), 130-133.
- McCulloch, S., et al., "Development of a fibre optic micro-optrode for intracellular pH measurements", IEE Proceedings, Optoelectronics, 144(3), (Jun. 1997), 162-167.
- Mintz, Gary S., et al., "American College of Cardiology Clinical Expert Consensus Document on Standards for Acquisition, Measurement and Reporting of Intravascular Ultrasound Studies (IVUS)", Journal of the American College of Cardiology, 37(5), (Apr. 2001), 1478-1492.
- Moreira, P. J., et al., "Dynamic Range Enhancement in Fiber Bragg Grating Sensors using a Multimode Laser Diode", IEEE Photonics Technology Letters, vol. 11, No. 6, (Jun. 1999), 3 pgs.
- O'Donnell, M., "New Imaging Technologies for Ultrasonography", J Med Ultrason., 27(4), Presented at the 73rd Japan Society of Ultrasound in Medicine Meeting, Invited Paper, (2000), 356-357.
- O'Donnell, M., et al., "Optoacoustics: high frequency ultrasonic array imaging", Proceedings of 17th International Congress on Acoustics, vol. IV, Biomedicine, Acoustics in Medicine, Invited Presentation at the 17th Intl Congress on Acoustics, Rome, (Sep. 27, 2001), 2-3.
- Othonos, A., et al., "In Section 7.9 Bragg Gating Fiber Laser Sensors from Fiber Bragg Gratings: fundamentals and applications in telecommunications and sensing", Artech House, Inc., (1999), 355-367.
- Pepine, Carl J., et al., "Improving Diagnostic and Therapeutic Outcomes Through Advanced Intravascular Imaging", Vascular Technologies, Inc., (1989), 3 pgs.
- Scully, P. J., "UV Laser Photo-induced Refractive Index Changes in Poly-Methyl-MethAcrylate and Plastic Optical Fibres for Application as Sensors and Devices", Central Laser Facility Annual Report, 1999/2000, (1999-2000), 145-147.
- Spisar, M., et al., "Stabilized, Resonant Optoacoustic Array Detectors for Medical Imaging", Proceedings of the World Congress on Ultrasonics, Paris, France, (Sep. 7-10, 2003), 25-28.
- Stefanadis, Christodoulos, et al., "Identification and Stabilization of Vulnerable Atherosclerotic Plaques: The Role of Coronary Tomography and External Heat Delivery", [online]. Retrieved from the Internet <URL: <http://www.indianheartjournal.org/Jan-Feb-2001/identification/identification/htm>>, (2001), 10 pgs.
- Surowiec, J., et al., "A Novel Miniature Optical Fibre Probe for MHz Frequency Ultrasound", Proceedings, IEEE Ultrasonics Symposium, vol. 2, (Nov. 3-6, 1996, San Antonio, TX), (1996), 1051-1054.
- Takahashi, N., et al., "Underwater Acoustic Sensor with Fiber Bragg Grating", Optical Review, 4(6), (1997), 691-694.
- Tanaka, S., et al., "Fibre optic spectral polarimetry for sensing multiple stress-loaded locations along a length of fibre", IEE Proceedings, Optoelectronics, 144(3), (Jun. 1997), 176-182.

(56)

References Cited

OTHER PUBLICATIONS

Wahle, Andreas, et al., "Accurate Visualization and Quantification of Coronary Vasculature by 3-D/4-D Fusion from Biplane Angiography and Intravascular Ultrasound", In: *Biomonitoring and Endoscopy Technologies*; I. Gannot et al., eds, (Jul. 5-6, 2000), 144-155.

Yoshino, T., et al., "Spiral fibre microbend sensors", *IEEE Proceedings, Optoelectronics*, 144(3), (Jun. 1997), 145-150.

"U.S. Appl. No. 12/020,736, Notice of Allowance mailed Aug. 29, 2014", 9 pgs.

"U.S. Appl. No. 14/053,421, Notice of Allowance mailed Nov. 24, 2014", 8 pgs.

"U.S. Appl. No. 14/053,421, Response filed Sep. 22, 2014 to Non Final Office Action mailed Jun. 20, 2014", 15 pgs.

"U.S. Appl. No. 14/490,464, Preliminary Amendment filed Sep. 19, 2014", 6 pgs.

Siebes, M., et al., "Single-Wire Pressure and Flow Velocity Measurement to Quantify Coronary Stenosis Hemodynamics and Effects of Percutaneous Interventions", *Circulation*, 109, (2004), 756-762.

"U.S. Appl. No. 14/053,421, Notice of Allowance mailed Mar. 11, 2015", 5 pgs.

"U.S. Appl. No. 14/490,464, Notice of Allowance mailed Mar. 17, 2015", 12 pgs.

"U.S. Appl. No. 14/577,980, Preliminary Amendment filed Mar. 25, 2015", 4 pgs.

U.S. Appl. No. 14/796,767, Non-Final Office Action mailed Sep. 29, 2015, 9 pgs.

U.S. Appl. No. 14/836,705, Preliminary Amendment filed Sep. 30, 2015, 5 pgs.

* cited by examiner

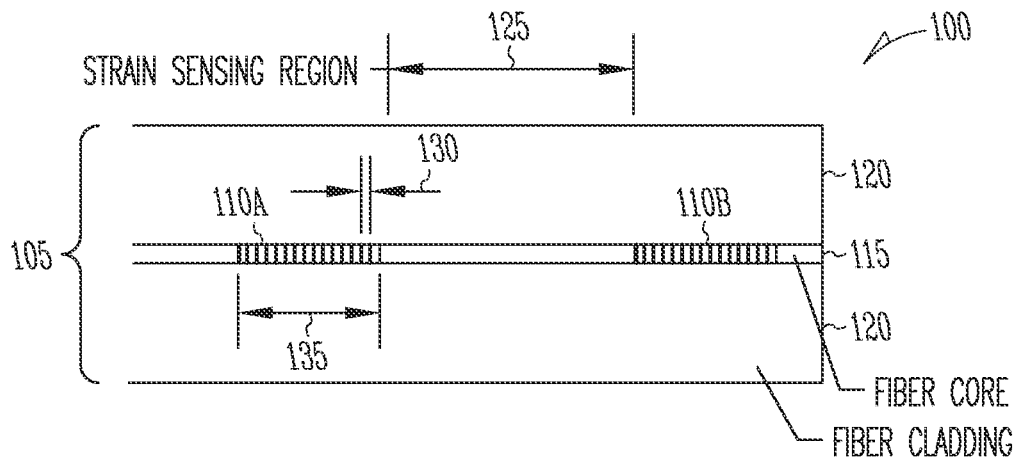


Fig. 1

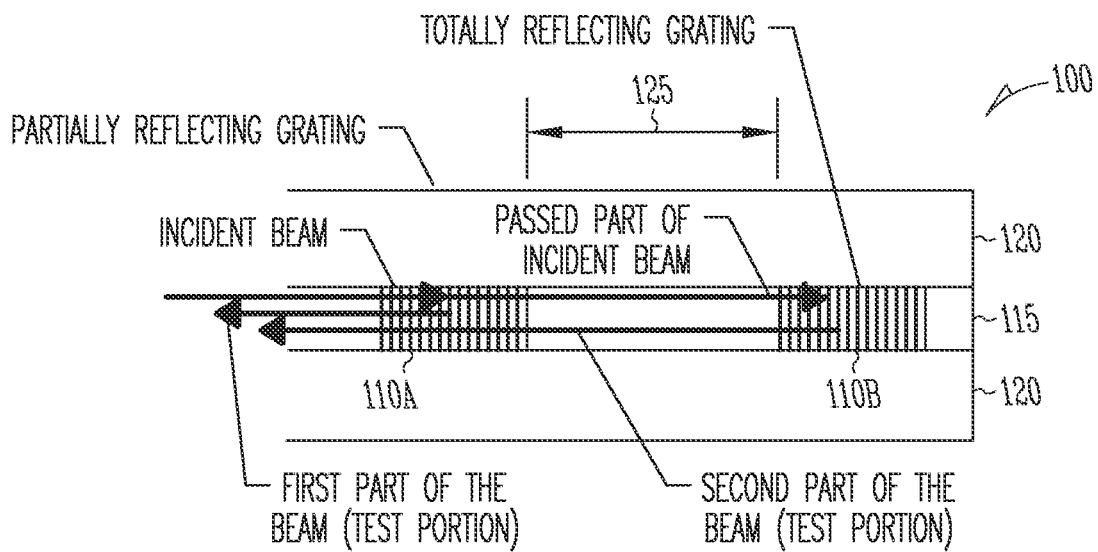


Fig. 2

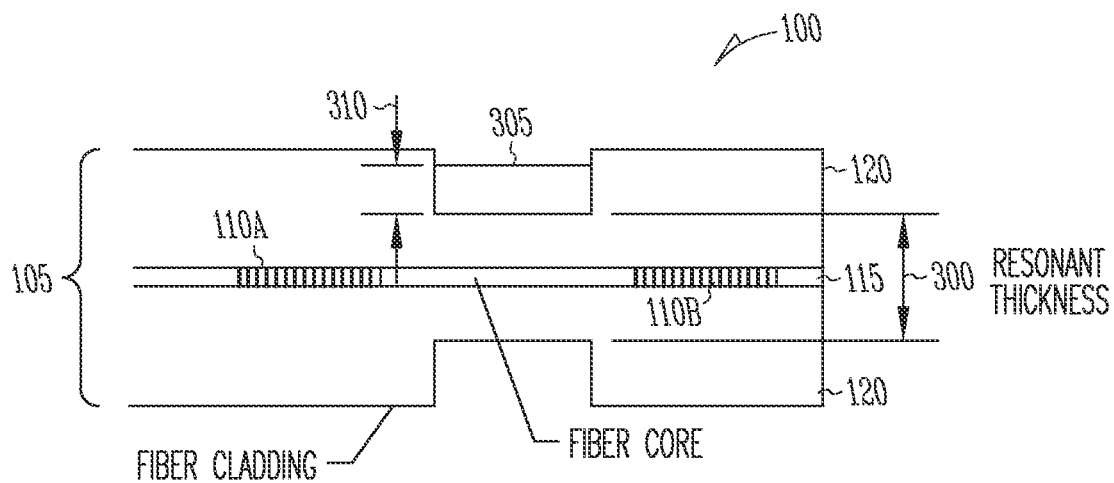


Fig. 3A

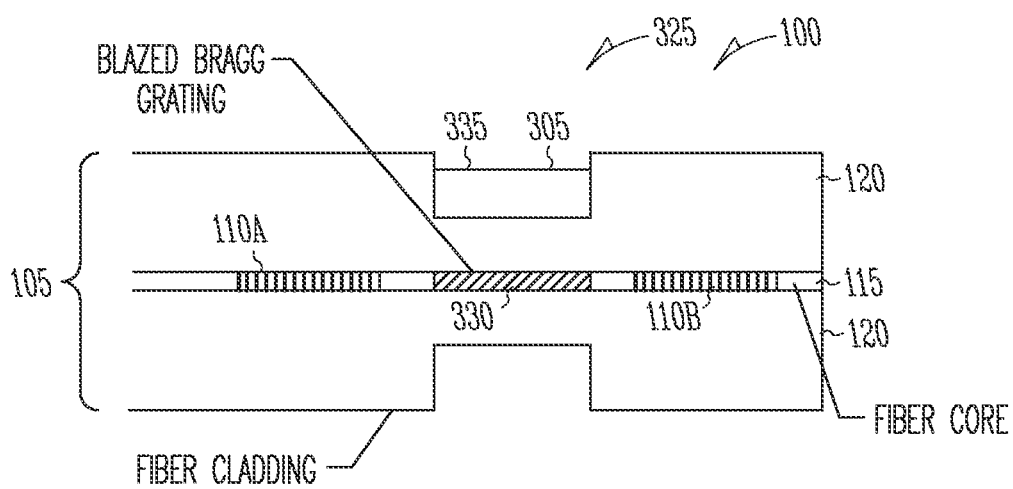


Fig. 3B

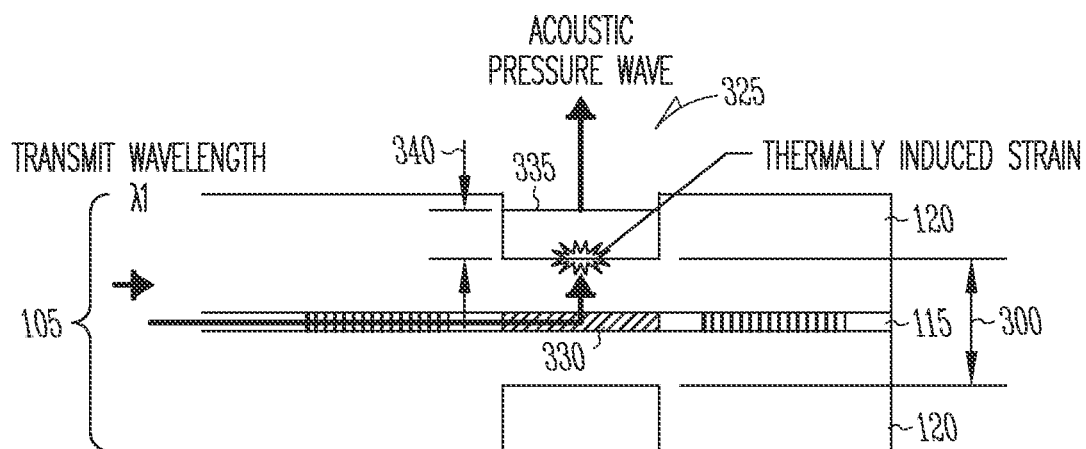


Fig. 4

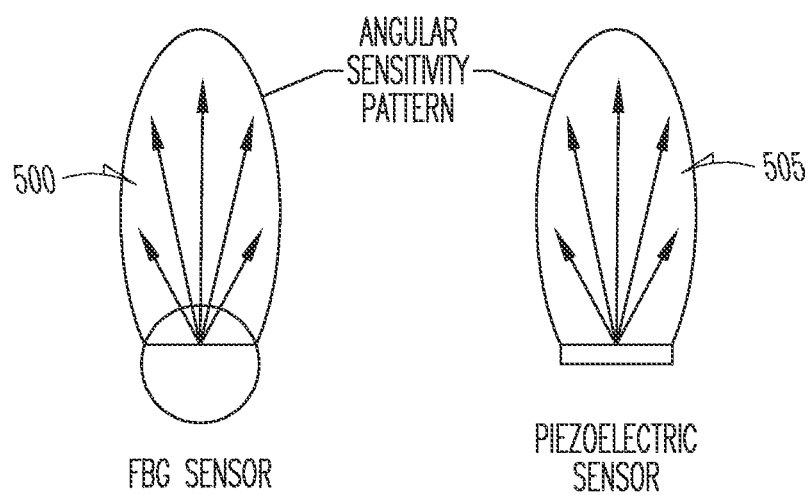


Fig. 5

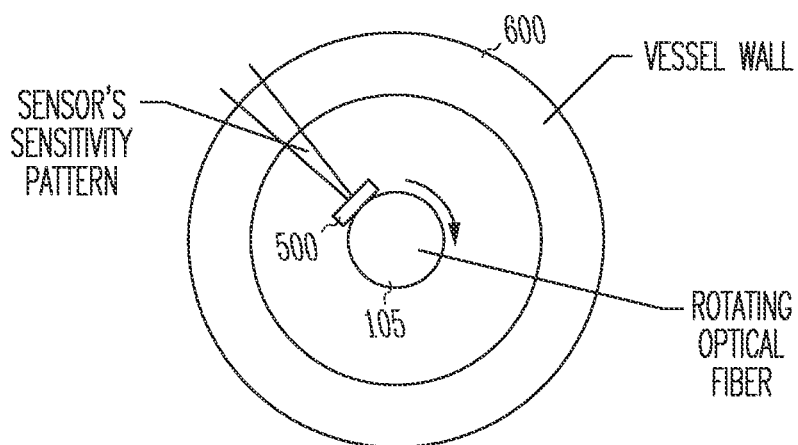


Fig. 6

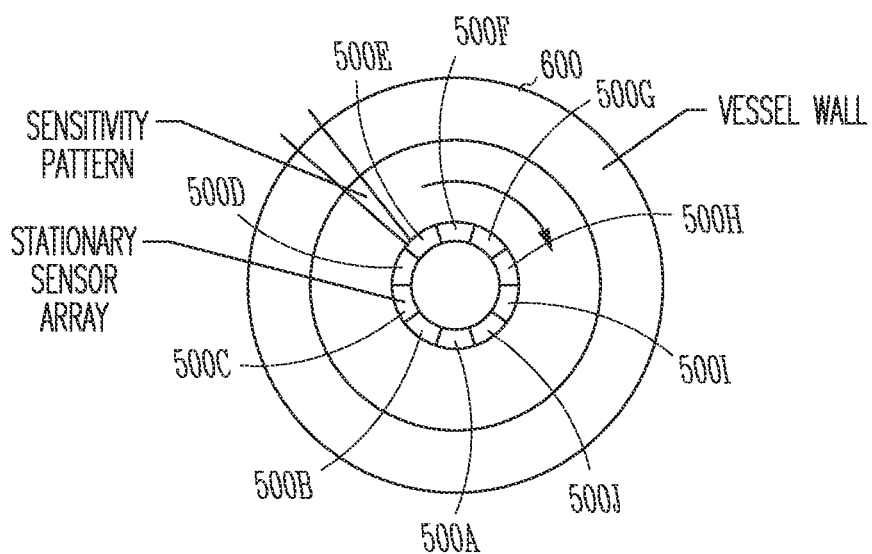


Fig. 7

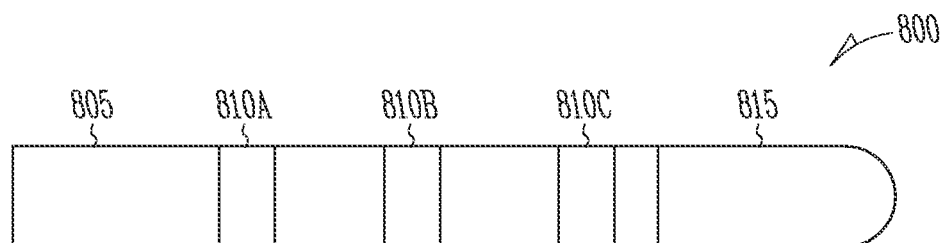


Fig. 8

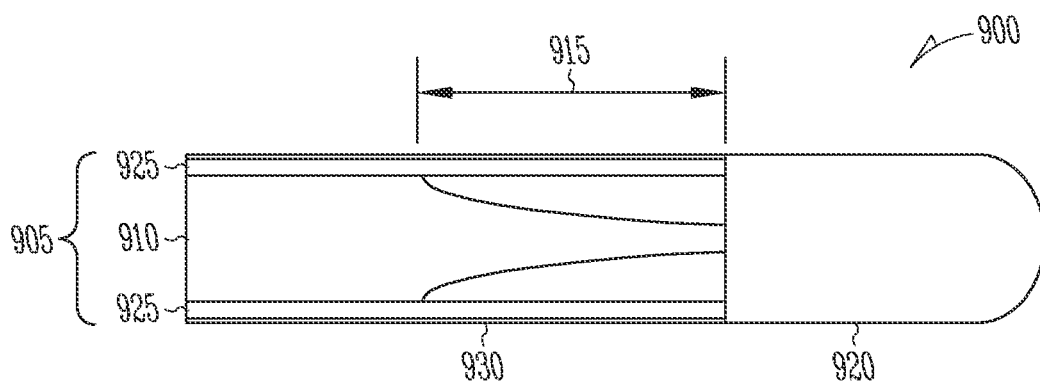


Fig. 9

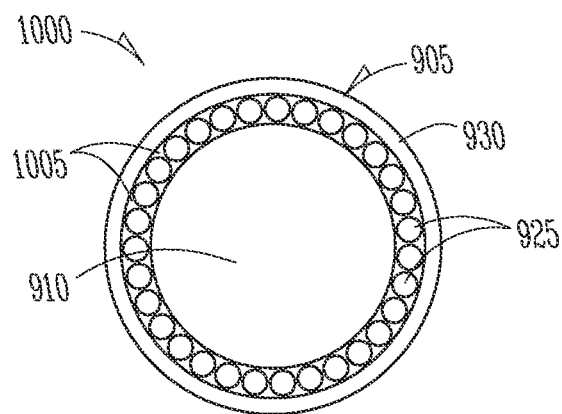


Fig. 10

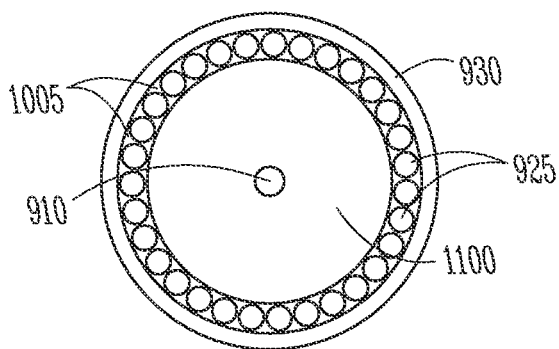


Fig. 11

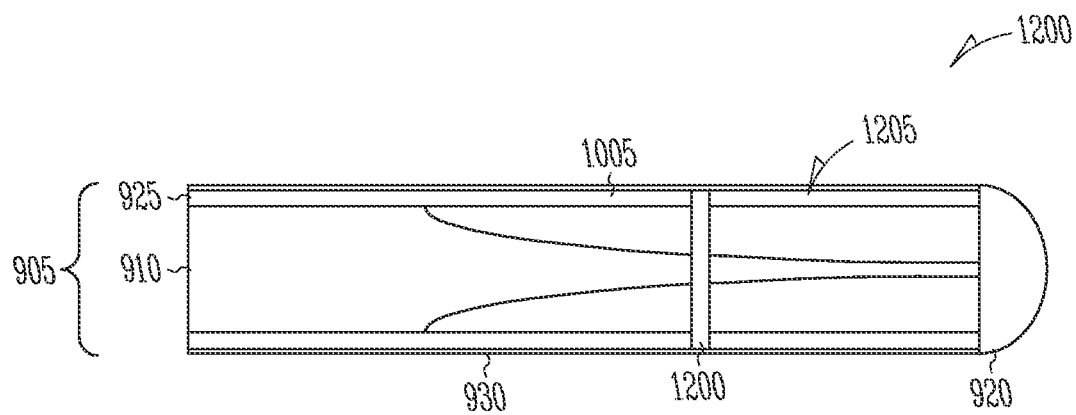


Fig. 12

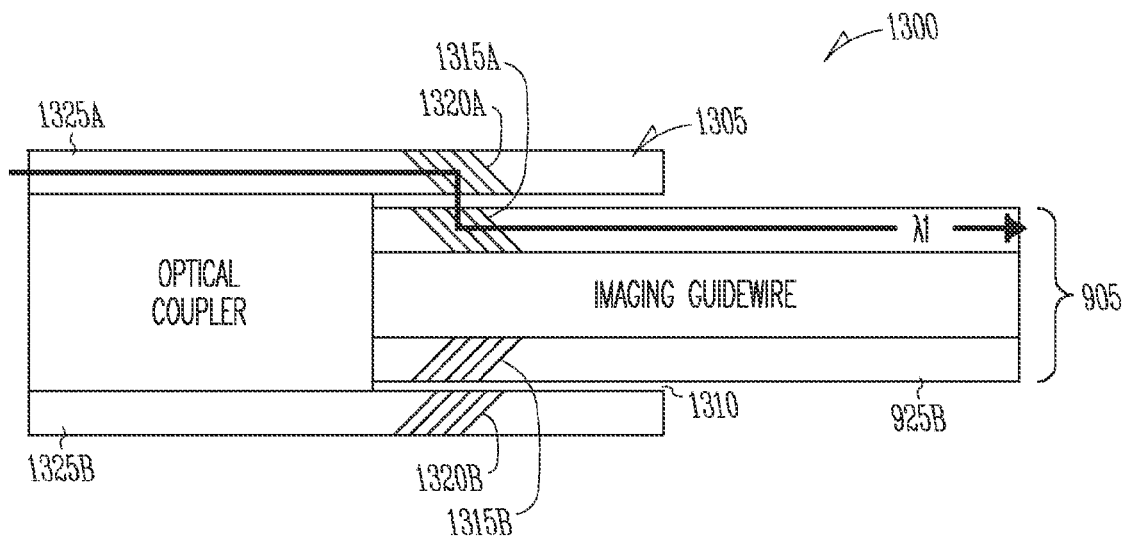


Fig. 13A

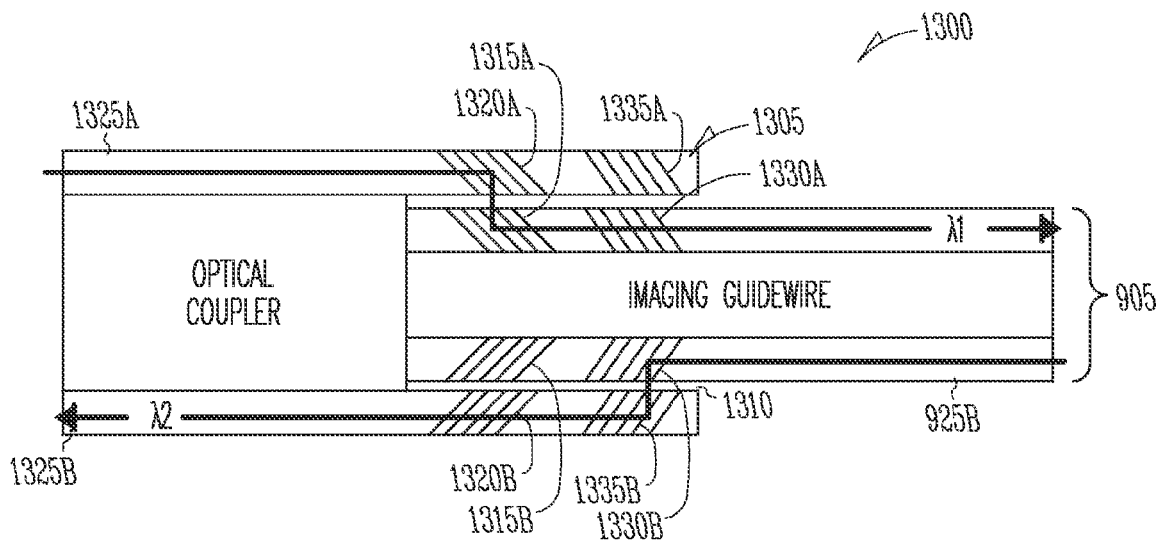
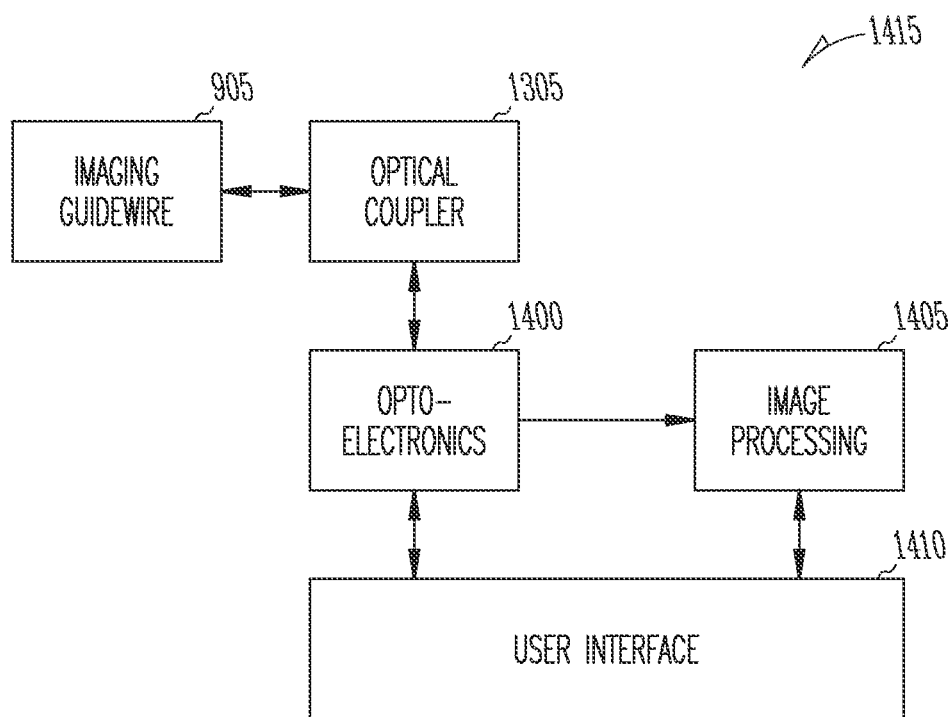
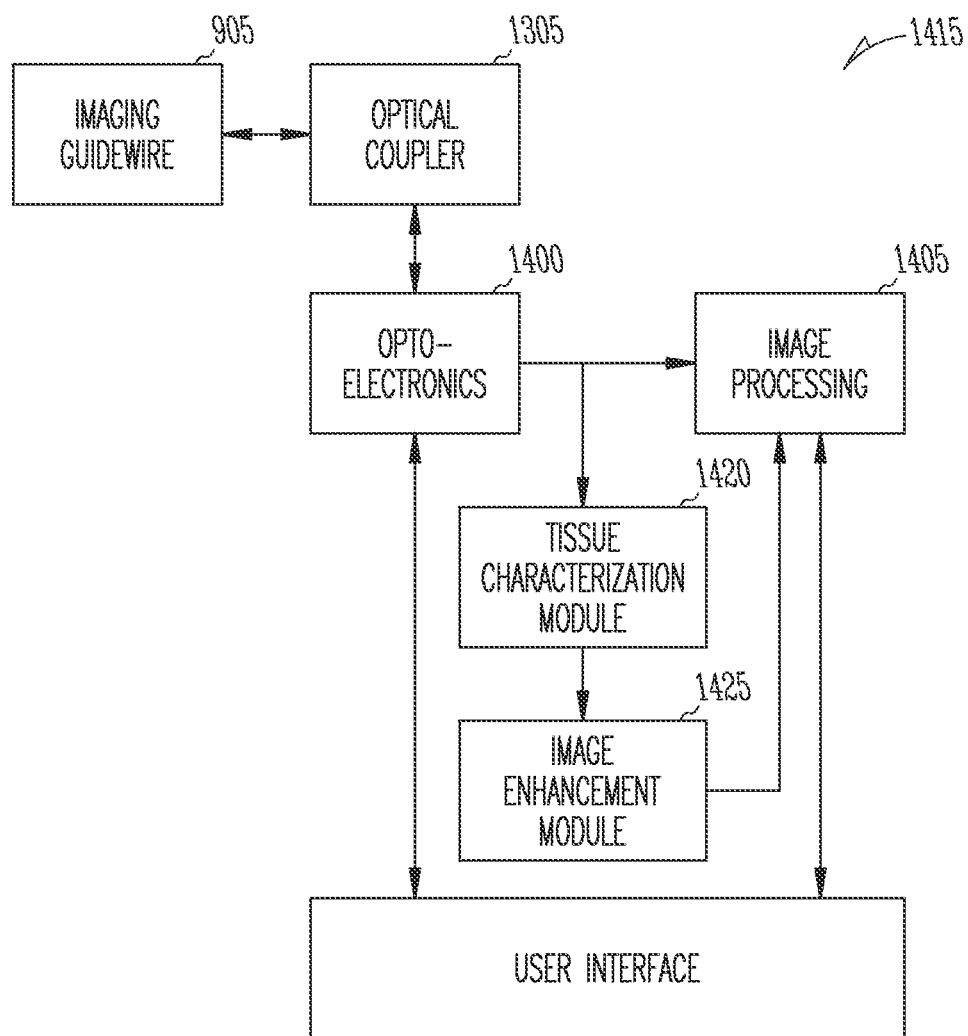


Fig. 13B

*Fig. 14A*

*Fig. 14B*

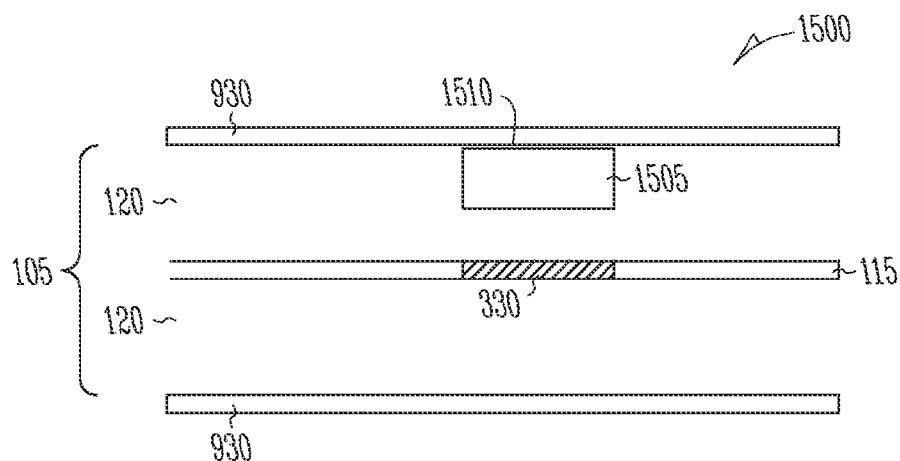


Fig. 15

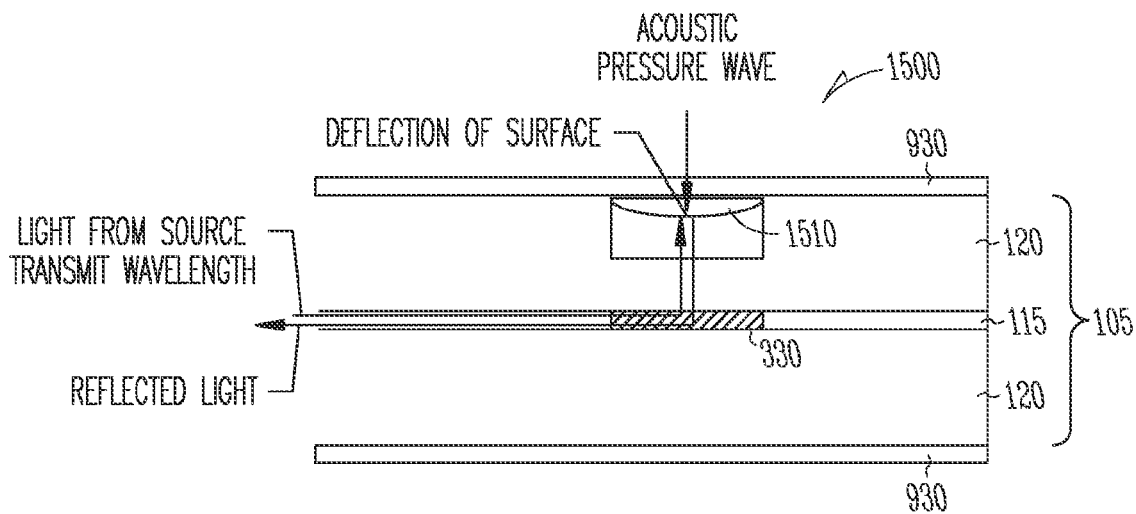


Fig. 16

SYSTEMS AND METHODS FOR MINIMALLY-INVASIVE OPTICAL-ACOUSTIC IMAGING

RELATED APPLICATION

This application is a continuation of U.S. patent application Ser. No. 13/779,985, filed on Feb. 28, 2013, issuing on May 20, 2014 as U.S. Pat. No. 8,731,340; which application is a continuation of U.S. patent application Ser. No. 13/285,551, filed on Oct. 31, 2011, now U.S. Pat. No. 8,391,652, issued on Mar. 5, 2013; which application is a continuation of U.S. patent application Ser. No. 12/701,228, filed on Feb. 5, 2010, now U.S. Pat. No. 8,059,923, issued on Nov. 15, 2011; which application is a continuation of U.S. patent application Ser. No. 12/263,978, filed on Nov. 3, 2008, now U.S. Pat. No. 7,660,492, issued on Feb. 9, 2010; which application is a continuation of U.S. patent application Ser. No. 11/674,568, filed on Feb. 13, 2007, now U.S. Pat. No. 7,447,388, issued on Nov. 4, 2008; which application is a continuation of U.S. patent application Ser. No. 10/266,082, filed on Oct. 7, 2002, now U.S. Pat. No. 7,245,789, issued on Jul. 17, 2007; all of which are incorporated herein by reference in their entireties.

FIELD OF THE INVENTION

This document relates generally to imaging, and particularly, but not by way of limitation, to systems and methods for minimally-invasive optical-acoustic imaging.

BACKGROUND

Vardi et al. PCT Patent Application No. PCT/US99/04913, which published on 18 Nov. 1999 as International Publication No. WO 99/58059, discusses an optical-acoustic imaging device. In that document, which is incorporated herein by reference in its entirety, including its disclosure of a guidewire optical-acoustic imaging device, an elongated imaging guidewire includes an elongated optical fiber driven by light from a laser. A distal end of the guidewire includes a polyvinylidene fluoride (PVDF) film piezoelectric ultrasound transducer. The ultrasound transducer transmits ultrasound to an imaging region of interest about the distal tip of the guidewire, and also receives the returned ultrasound energy. The received ultrasound energy deforms a Fiber Bragg Grating (FBG or "Bragg grating") at the distal end of the guidewire, which, in turn, modulates the optical signal through the optical fiber. Imaging information about the region of interest is then obtained at the proximal end of the guidewire from the modulated optical signal.

Among other things, the present applicant has recognized that a piezoelectric ultrasound transducer may be difficult to integrate with a minimally-invasive guidewire because of electrical signal losses in ultrafine electrical conductors extending longitudinally through the guidewire assembly. Moreover, the present applicant has recognized that the field of view of the Vardi et al. device may be limited by the size of apertures around the PVDF ultrasound transducers and/or the spacing between FBGs. Furthermore, the present applicant has recognized that the sensitivity of the Vardi et al. device may limit its usefulness in an imaging application. For these and other reasons, the present applicant has recognized that there is an unmet need in the art for improved systems and methods for performing optical-acoustic imaging.

BRIEF DESCRIPTION OF THE DRAWINGS

In the drawings, which are not necessarily drawn to scale, like numerals describe substantially similar components

throughout the several views. Like numerals having different letter suffixes represent different instances of substantially similar components. The drawings illustrate generally, by way of example, but not by way of limitation, various embodiments discussed in the present document.

FIG. 1 is a cross-sectional side view illustrating generally, by way of example, but not by way of limitation, one example of an FBG strain sensor in an optical fiber.

FIG. 2 is a cross-sectional side view illustrating generally, by way of example, but not by way of limitation, an example of an FBG grating interferometer sensor.

FIG. 3A is a cross-sectional schematic diagram illustrating generally one example of a distal portion of an imaging guidewire that combines an acousto-optic FBG sensor with a photoacoustic transducer.

FIG. 3B is a cross-sectional schematic diagram illustrating generally one example of a distal portion of an imaging guidewire that combines an acousto-optic FBG sensor with a photoacoustic transducer.

FIG. 4 is a cross-sectional schematic diagram illustrating generally one example of the operation of a blazed grating FBG photoacoustic transducer.

FIG. 5 is a schematic diagram comparing an expected angular sensitivity pattern of an exemplary blazed FBG optical-to-acoustic and acoustic-to-optical combined device to that of a piezoelectric sensor.

FIG. 6 is a schematic diagram illustrating generally one technique of generating an image by rotating the blazed FBG optical-to-acoustic and acoustic-to-optical combined transducer and displaying the resultant series of radial image lines to create a radial image.

FIG. 7 is a schematic diagram that illustrates generally one such phased array example, in which the signal to/from each array transducer is combined with the signals from the other transducers to synthesize a radial image line.

FIG. 8 is a schematic diagram that illustrates generally an example of a side view of a distal portion of a guidewire.

FIG. 9 is a schematic diagram that illustrates generally one example of a cross-sectional side view of a distal portion of a guidewire.

FIG. 10 is a schematic diagram that illustrates generally one example of a cross-sectional end view of a proximal portion of a guidewire.

FIG. 11 is a schematic diagram that illustrates generally one example of a cross-sectional end view of a distal portion of a guidewire.

FIG. 12 is a schematic diagram that illustrates generally one example of cross-sectional side view of a distal portion of a guidewire.

FIG. 13A is a cross-sectional schematic diagram illustrating generally one example of a proximal portion of a guidewire, which is communicatively coupled to an instrumentation/control interface via an optical coupler.

FIG. 13B is a cross-sectional schematic diagram illustrating generally a further example of a proximal portion of a guidewire that is communicatively coupled to an instrumentation/control interface using an optical coupler.

FIG. 14A is a block diagram illustrating generally one example of the imaging guidewire and associated interface components.

FIG. 14B is a block diagram illustrating generally another example of the imaging guidewire and associated interface components, including tissue characterization and image enhancement modules.

FIG. 15 is a cross-sectional schematic diagram illustrating generally one example of an alternate acoustic-to-optical transducer.

FIG. 16 is a cross-sectional schematic diagram illustrating generally one example of operation of the acoustic-to-optical transducer of FIG. 15.

DETAILED DESCRIPTION

In the following detailed description, reference is made to the accompanying drawings that form a part hereof, and in which is shown by way of illustration specific embodiments in which the invention may be practiced. These embodiments are described in sufficient detail to enable those skilled in the art to practice the invention, and it is to be understood that the embodiments may be combined, or that other embodiments may be utilized and that structural, logical and electrical changes may be made without departing from the scope of the present invention. The following detailed description is, therefore, not to be taken in a limiting sense, and the scope of the present invention is defined by the appended claims and their equivalents.

In this document, the terms “a” or “an” are used, as is common in patent documents, to include one or more than one. Furthermore, all publications, patents, and patent documents referred to in this document are incorporated by reference herein in their entirety, as though individually incorporated by reference. In the event of inconsistent usages between this documents and those documents so incorporated by reference, the usage in the incorporated reference(s) should be considered supplementary to that of this document; for irreconcilable inconsistencies, the usage in this document controls. In this document, the term “minimally-invasive” refers to techniques that are less invasive than conventional surgery; the term “minimally-invasive” is not intended to be restricted to the least-invasive technique possible.

1. EXAMPLES OF FIBER BRAGG GRATING ACOUSTO-OPTIC SENSORS

FIG. 1 is a cross-sectional side view illustrating generally, by way of example, but not by way of limitation, one example of a strain-detecting FBG sensor 100 in an optical fiber 105. FBG sensor 100 senses acoustic energy received from a nearby area to be imaged, and transduces the received acoustic energy into an optical signal within optical fiber 105. In the example of FIG. 1, FBG sensor 100 includes Bragg gratings 110A-B in an optical fiber core 115 surrounded by an optical fiber cladding 120. Bragg gratings 110A-B are separated by a strain sensing region 125, which, in one example, is about a millimeter in length. This example senses strain by detecting an “optical displacement” between these gratings 110A-B.

A fiber Bragg grating can be conceptualized as a periodic change in the optical index (which is inversely proportional to the speed of light in the material) of a portion of the optical fiber core 115. Light of a specific wavelength traveling down such a portion of core 115 will be reflected; the period (distance) 130 of the change in the optical index determines the particular wavelength of light that will be reflected. The degree of index change and the length 135 of the grating determine the ratio of light reflected to that transmitted through the grating.

FIG. 2 is a cross-sectional side view illustrating generally, by way of example, but not by way of limitation, an operative example of an interferometric FBG sensor 100. The example of FIG. 2 includes two FBGs 110A-B. FBG 110A is partially reflective at a specific wavelength of light passing through fiber core 115. FBG 110B is substantially fully reflective of such light. This interferometric arrangement of FBGs 110A-B is capable of discerning the “optical distance”

between FBGs 110A-B with extreme sensitivity. The “optical distance” is a function of the velocity of light in the material of fiber core 115 as well as the length 125 between FBGs 110A-B. Thus, a change in the velocity of light can induce a change in optical distance even though the physical distance 125 between FBGs 110A-B has not changed.

An interferometer such as FBG sensor 100 can be conceptualized as a device that measures the interference between two paths taken by an optical beam. A partially reflecting FBG 110A (or a partially reflecting mirror) is used to split the incident beam of light into two parts. In an interferometer, one part of the beam travels along a path that is kept constant (i.e., a control path) and the other part travels a path where some change is to be monitored (i.e., a test path). Using partially reflecting FBG 110A (or a partially reflecting mirror, either of which may alternatively be in addition to FBG 110A), the two parts of the beam are combined. If the two paths are identical, the parts combine to form the original beam. If the paths are different, the two parts will add or subtract from each other. This addition or subtraction is known as interference. A complete subtraction is called a null and occurs at a precise wavelength of light for a given difference in paths. Measuring the wavelength where this null occurs yields an indication of the difference in optical paths between the two beams. In such a manner, an interferometer such as FBG sensor 100 senses small changes in distance, such as a change in the optical distance 125 between FBGs 110A-B resulting from received ultrasound or other received acoustic energy.

In one example, such as illustrated in FIG. 2, the interferometric FBG sensor 100 causes the interference between that portion of the optical beam that is reflected off the first (partially reflective) FBG 110A with that reflected from the second (substantially fully reflective) FBG 110B. The wavelength of light where an interferometric null will occur is very sensitive to the “optical distance” 125 between the two FBGs 110A-B. This interferometric FBG sensor 100 of FIG. 2 has another very practical advantage. In this example, the two optical paths along the fiber core 115 are the same, except for the sensing region between FBGs 110A-B. This shared path ensures that any optical changes in the shared portion of optical fiber 105 will have substantially no effect upon the interferometric signal; only the change in the sensing region between FBGs 125 is sensed.

2. EXAMPLES OF DEVICES AND METHODS IMPROVING FBG RECEPTION

In one example, an FBG sensor 100 senses strain generated by ultrasound or other acoustic energy received from a nearby imaging region to be visualized and, in response, modulates an optical signal in an optical fiber. Increasing the sensitivity of the FBG sensor 100 provides improved imaging. A first example of increasing sensitivity is to increase the amount of strain induced in the FBG sensor 100 for a given dynamic pressure provided by the acoustic energy. A second example is to increase the modulation of the optical signal for a given change in strain of the FBG sensor 100.

One technique of increasing the strain induced in the FBG sensor 100 is to design the physical attributes of the FBG sensor 100 to increase the degree of strain for a given externally applied acoustic field. In one such example, increased strain is obtained by using material that has a high degree of strain for a given stress. Calculations indicate that over two orders of magnitude increase in strain vs. stress (also referred to as “compliance”) is obtained by using an optical grade plastic, rather than glass, in the fiber core 115 of the FBG

5

sensor **100**. One example of a suitable optical grade plastic used in fiber core **115** is poly-methyl-methacrylate (PMMA).

In a second example, the FBG sensor **100** is shaped so as to increase the strain for a given applied acoustic pressure field. FIG. **3A** is a cross-sectional schematic diagram illustrating one such example in which the FBG sensor **100** is shaped such that it mechanically resonates at the frequency of the acoustic energy received from the nearby imaging region, thereby resulting in increased strain. In the example of FIG. **3A**, all or a portion of the strain sensing region between FBGs **110A-B** is selected to provide a resonant thickness **300** that promotes mechanical resonance of the received acoustic energy, thereby increasing the resulting strain sensed by FBG sensor **100**. In one example, such as illustrated in FIG. **3A**, this is accomplished by grinding or otherwise removing a portion of fiber cladding **120**, such that the remaining thickness of fiber core **115** and/or fiber cladding **120** between opposing planar (or other) surfaces is selected to mechanically resonate at the frequency of the acoustic energy received from the nearby imaging region.

In one example, for a particular material, mechanical resonance is obtained by making the resonant thickness **300** of the strain sensing region substantially the same thickness as $\frac{1}{2}$ the acoustic wavelength (or an odd integer multiple thereof) in the material(s) of FBG sensor **100** at the acoustic center frequency of the desired acoustic frequency band received from the imaging region. In other examples, such as for other materials, the resonant thickness **300** is selected to match a different proportion of the acoustic wavelength that obtains the desired mechanical resonance for that material. Calculations indicate that obtaining such mechanical resonance will increase the strain sensitivity by about an order of magnitude over that of a sensor that is not constructed to obtain such mechanical resonance.

In a third example, a special coating **305** is applied to the FBG sensor **100** to increase the acoustic pressure as seen by the FBG sensor **100** over a band of acoustic frequencies, thereby improving its sensitivity over that band. The difference between the mechanical characteristics of water (or tissue and/or blood, which is mostly comprised of water) and glass material of the optical fiber **105** carrying the FBG sensor **100** is typically so significant that only a small amount of acoustic energy “enters” the FBG sensor **100** and thereby causes strain; the remaining energy is reflected back into the biological or other material being imaged. For a particular range of acoustic frequencies, one or more coatings **305** of specific thickness **310** and/or mechanical properties (e.g., the particular mechanical impedance) of the coating material can dramatically reduce such attenuation due to the different mechanical characteristics. One example uses quarter wave matching, providing a coating **305** of a thickness **310** that is approximately equal to one quarter of the acoustic signal wavelength received from the region being imaged. Using such matching, the sensitivity of the FBG sensor **100**, over a given band of acoustic frequencies of interest, is expected to increase by about an order of magnitude.

In one example, using the above-discussed quarter wave matching and sensor shaping techniques, the sensitivity of the FBG sensor **100** approaches that of a piezoelectric transducer. Additionally using optical grade plastic for fiber core **115**, in conjunction with one or the other of these techniques, will further increase the sensitivity of the FBG sensor **100**.

Additionally (or alternatively) to the above techniques of increasing the strain of the FBG sensor **100** for a particular level of acoustic energy, in one example, the optical sensitivity of the FBG sensor **100** to strain is increased, thereby increasing the sensitivity of the FBG sensor **100** to an acoustic

6

field. In one example, this is accomplished by improved techniques of optical wavelength discrimination, such as by using a fiber-based Mach-Zehnder interferometer or by construction of improved optical fiber geometries that increase the optical sensitivity of the FBG sensor **100** to strain.

3. EXAMPLES OF SYSTEMS AND METHODS IMPROVING ULTRASOUND TRANSMISSION

The present applicant has recognized that while it may be possible to implement an imaging guidewire that transmits ultrasound using a piezoelectric transducer, such a design may involve a trade-off. If the piezoelectric transducer radiates the ultrasonic energy in a broad radial pattern, imaging quality may be degraded. Conversely, using smaller piezoelectric transducers to transmit ultrasound may require significant electrical voltages in a guidewire in order to achieve needed acoustic transmit energy. This is because a smaller transducer has a higher electrical impedance, needing a higher voltage to achieve the same acoustic power. Such a guidewire must also use materials of sufficient dielectric properties to ensure patient safety. Moreover, adding electrically conducting wires to a guidewire assembly complicates its manufacture.

However, optical energy can be converted to acoustic energy. In one example, therefore, such problems are overcome using an optical-to-acoustic transmitter, which, in one example, is integrated with an acoustic-to-optical receiver such as FBG sensor **100**.

FIG. **3B** is a cross-sectional schematic diagram illustrating generally one example of a distal portion of an imaging guidewire that combines an acousto-optic FBG sensor **100** with an photoacoustic transducer **325**. In this example, photoacoustic transducer **325** includes a blazed Bragg grating **330**. In the illustrative example of FIG. **3B**, blazed Bragg grating **330** is implemented in the strain sensitive region of the FBG sensor **100**, between FBGs **110A-B**, however, this is not a requirement. Unlike an unblazed Bragg grating, which typically includes impressed index changes that are substantially perpendicular to the longitudinal axis of the fiber core **115** of the optical fiber **105**, the blazed Bragg grating **330** includes obliquely impressed index changes that are at a nonperpendicular angle to the longitudinal axis of the optical fiber **105**.

A standard unblazed FBG partially or substantially fully reflects optical energy of a specific wavelength traveling down the axis of the fiber core **115** of optical fiber **105** back up the same axis. Blazed FBG **330** reflects this optical energy away from the longitudinal axis of the optical fiber **105**. For a particular combination of blaze angle and optical wavelength, the optical energy will leave blazed FBG **330** substantially normal (i.e., perpendicular) to the longitudinal axis of the optical fiber **105**. In the illustrative example of FIG. **3B**, an optically absorptive photoacoustic material **335** (also referred to as a “photoacoustic” material) is placed on the surface of optical fiber **105**. The optically absorptive photoacoustic material **335** is positioned, with respect to the blazed grating **330**, so as to receive the optical energy leaving the blazed grating. The received optical energy is converted in the optically absorptive material **335** to heat that expands the optically absorptive photoacoustic material **335**. The optically absorptive photoacoustic material **335** is selected to expand and contract quickly enough to create and transmit an ultrasound or other acoustic wave that is used for acoustic imaging of the region of interest about the distal tip (or other desired portion) of the imaging guidewire. In one example, the optically absorptive photoacoustic material **335** is the same material as the acoustic matching material **305** discussed above.

FIG. 4 is a cross-sectional schematic diagram illustrating generally one example of the operation of photoacoustic transducer 325 using a blazed Bragg grating 330. Optical energy of a specific wavelength, λ_1 , travels down the fiber core 115 of optical fiber 105 and is reflected out of the optical fiber 105 by blazed grating 330. The outwardly reflected optical energy impinges on the photoacoustic material 335. The photoacoustic material 335 then generates a responsive acoustic impulse that radiates away from the photoacoustic material 335 toward nearby biological or other material to be imaged. Acoustic energy of a specific frequency is generated by optically irradiating the photoacoustic material 335 at a pulse rate equal to the desired acoustic frequency.

In another example, the photoacoustic material 335 has a thickness 340 (in the direction in which optical energy is received from blazed Bragg grating 330) that is selected to increase the efficiency of emission of acoustic energy. In one example, thickness 340 is selected to be about $\frac{1}{4}$ the acoustic wavelength of the material at the desired acoustic transmission/reception frequency. This improves the generation of acoustic energy by the photoacoustic material.

In yet a further example, the photoacoustic material is of a thickness 300 that is about $\frac{1}{4}$ the acoustic wavelength of the material at the desired acoustic transmission/reception frequency, and the corresponding glass-based optical fiber sensing region resonant thickness 300 is about $\frac{1}{2}$ the acoustic wavelength of that material at the desired acoustic transmission/reception frequency. This further improves the generation of acoustic energy by the photoacoustic material and reception of the acoustic energy by the optical fiber sensing region.

In one example of operation, light reflected from the blazed grating excites the photoacoustic material in such a way that the optical energy is efficiently converted to substantially the same acoustic frequency for which the FBG sensor is designed. The blazed FBG and photoacoustic material, in conjunction with the aforementioned FBG sensor, provide both a transmit transducer and a receive sensor, which are harmonized to create an efficient unified optical-to-acoustic-to-optical transmit/receive device. In one example, the optical wavelength for sensing is different from that used for transmission. In a further example, the optical transmit/receive frequencies are sufficiently different that the reception is not adversely affected by the transmission, and vice-versa.

FIG. 5 is a schematic diagram comparing an expected angular sensitivity pattern of an exemplary blazed FBG optical-to-acoustic and acoustic-to-optical combined device 500 to that of a piezoelectric transducer 505. As seen in the example of FIG. 5, the optical-to-acoustic-to-optical sensor assembly 500 is expected to be capable of operating over a specific angular range that is substantially similar to that of the piezoelectric transducer 505 of similar dimensions. Therefore, in one example, the blazed optical-to-acoustic and acoustic-to-optical combined device 500 is capable of using conventional intravascular ultrasound ("IVUS") techniques.

FIG. 6 is a schematic diagram illustrating generally one technique of generating an image of a vessel wall 600 by rotating the blazed FBG optical-to-acoustic and acoustic-to-optical combined transducer 500 and displaying the resultant series of radial image lines to create a radial image. In another example, phased array images are created using a substantially stationary (i.e., non-rotating) set of multiple FBG sensors, such as FBG sensors 500A-J. FIG. 7 is a schematic diagram that illustrates generally one such phased array example, in which the signal to/from each array transducer 500A-J is combined with the signals from one or more other transducers 500A-J to synthesize a radial image line. In this

example, other image lines are similarly synthesized from the array signals, such as by using specific changes in the signal processing used to combine these signals.

4. EXAMPLES OF GUIDEWIRE DESIGN

FIG. 8 is a schematic diagram that illustrates generally an example of a side view of a distal portion 800 of an imaging guidewire 805 or other elongate catheter (in one example, the guidewire 805 is capable of being used for introducing and/or guiding a catheter or other medical instrument, e.g., over the guidewire 805). In this example, the distal portion 800 of the imaging guidewire 805 includes one or more imaging windows 810A, 810B, . . . , 810N located slightly or considerably proximal to a distal tip 815 of the guidewire 805. Each imaging window 810 includes one or more optical-to-acoustic transducers 325 and a corresponding one or more separate or integrated acoustic-to-optical FBG sensors 100. In one example, each imaging window 810 includes an array of blazed FBG optical-to-acoustic and acoustic-to-optical combined transducers 500 (such as illustrated in FIG. 7) located slightly proximal to distal tip 815 of guidewire 805 having mechanical properties that allow the guidewire 805 to be guided through a vascular or other lumen. In one example, the different imaging windows 810A, 810B, . . . , 810N are designed for different optical wavelengths, such that individual windows can be easily addressed by changing the optical wavelength being communicated through fiber core 115.

FIG. 9 is a schematic diagram that illustrates generally one example of a cross-sectional side view of a distal portion 900 of another guidewire 905. In this example, the guidewire 905 includes a solid metal or other core 910 that tapers down in diameter (e.g., from an outer diameter of about 0.011 inches) at a suitable distance 915 (e.g., about 50 cm) from the distal tip 920, to which the tapered core 910 is attached. In this example, optical fibers 925 are distributed around the outer circumference of the guidewire core 910, and attached to the distal tip 920. In this example, the optical fibers 925 are at least partially embedded in a polymer matrix or other binder material that bonds the optical fibers 925 to the guidewire core 910 and/or the distal tip 920. The binder material may also contribute to the torsion response of the resulting guidewire assembly 905. In one example, the optical fibers 925 and binder material is overcoated with a polymer or other coating 930, such as for providing abrasion resistance, optical fiber protection, and/or friction control. In this example, the composite structure of the distal region 900 of the guidewire 905 provides, among other things, flexibility and rotational stiffness, thereby allowing the guidewire 905 to be maneuvered to an imaging region of interest within a vascular or any other lumen.

FIG. 10 is a schematic diagram that illustrates generally one example of a cross-sectional end view of a proximal portion 1000 of guidewire 905, which includes guidewire core 910, optical fibers 925, binder material 1005, and outer coating 930. In this example, but not by way of limitation, the diameter of the core 910 is about $\frac{1}{1000}$ inch, the diameter of the optical fibers 925 is about $(1.25)/1000$ inch, and the optional outer coating 930 is about $(0.25)/1000$ inch thick.

FIG. 11 is a schematic diagram that illustrates generally one example of a cross-sectional end view of distal portion 900 of guidewire 905, e.g., adjacent to distal tip 920. In this example, but not by way of limitation, the diameter of core 910 has tapered down to about $(1.5)/1000$ inch, circumferentially surrounded by a void 1100 of about the same diameter (e.g., about $\frac{1}{1000}$ inch) as the core 910 near the proximal end

100 of the guidewire 905. In this example, the optical fibers 925 are circumferentially disposed in the binder material 1005 around the void 1100. Binder material 1005 provides structural support. Optical fibers 925 are optionally overlaid with the outer coating 930.

FIG. 12 is a schematic diagram that illustrates generally one example of a cross-sectional side view of a distal portion 900 of a guidewire 905. In this example, at least one metallic or other bulkhead 1200 is provided along the tapered portion of the guidewire core 910. The optical fibers 925 and binder 1005 are attached to a proximal side of the bulkhead 1200 near its circumferential perimeter. A distal side of the bulkhead 1200 is attached, near its circumferential perimeter, to a coil winding 1205 that extends further, in the distal direction, to a ball or other distal tip 920 of the guidewire 905.

5. EXAMPLES OF ACOUSTIC TRANSDUCER CONSTRUCTION

In one example, before the acoustic transducer(s) is fabricated, the guidewire 905 is assembled, such as by binding the optical fibers 925 to the core 910 and distal tip 920 or bulkhead 1200, and optionally coating the guidewire 905. The optoacoustic transducer(s) are then integrated into the guidewire assembly 905, such as by grinding one or more grooves in the guidewire at the locations of the optoacoustic transducer windows 810. In a further example, the depth of these groove(s) in the optical fiber(s) 925 defines the resonant structure(s) of the optoacoustic transducer(s).

After the optoacoustic transducer windows 810 have been defined, the FBGs added to one or more portions of the optical fiber 925 within such windows 810. In one example, the FBGs are created using an optical process in which the portion of the optical fiber 925 is exposed to a carefully controlled pattern of UV radiation that defines the Bragg gratings. Then, a photoacoustic material is deposited or otherwise added in the transducer windows 810 over respective Bragg gratings. One example of a suitable photoacoustic material is pigmented polydimethylsiloxane (PDMS), such as a mixture of PDMS, carbon black, and toluene. Thus, in this example, the FBGs are advantageously constructed after the major elements of the guidewire are mechanically assembled into the guidewire assembly 905.

6. EXAMPLES OF PROXIMAL END INTERFACE

In one example, the guidewire 905 allows for over-the-guidewire or other insertion of a catheter at the proximal end of the guidewire. Therefore, in such an example, the guidewire 905 (including any proximal end interface) has an outer diameter that is less than or equal to the inner diameter (e.g., 0.014 inches) of a catheter to allow the catheter with a similarly-sized inner diameter to travel over the guidewire 905. FIGS. 13A and 13B provide illustrative examples of an optical coupler that easily engages and disengages guidewire 905. Among other things, this facilitates over-the-guidewire catheter insertion, and viewing an imaging region either before, during, or after such a catheter is inserted over-the-guidewire.

FIG. 13A is a cross-sectional schematic diagram illustrating generally one example of a proximal portion 1300 of guidewire 905, which is communicatively coupled to an instrumentation/control interface via an optical coupler 1305. In this example, proximal portion 1300 of guidewire 905 is received within a receptacle 1310 portion of optical coupler 1305, and includes one or more blazed FBGs 1315 to couple light into and/or out of one or more respective optical fibers

925 of guidewire 905. Optical coupler 1305 includes one or more corresponding blazed FBGs 1320 to couple light into and/or out of one or more respective optical fibers 1325 of optical coupler 1305. In the example of FIG. 13A, the FBGs 1320 of optical coupler 1305 are located substantially adjacent to corresponding FBGs 1315 of guidewire 905 when guidewire 905 is engaged within receptacle 1310 of optical coupler 1305. Although FIG. 13A illustrates a multiple-fiber embodiment of guidewire 905, the illustrated techniques for coupling to an instrumentation/control interface are also applicable to a guidewire that includes a single optical fiber.

FIG. 13B is a cross-sectional schematic diagram illustrating generally a further example of a proximal portion 1300 of guidewire 905 that is communicatively coupled to an instrumentation/control interface using an optical coupler 1305. In the example of FIG. 13B, at least one optical fiber 925 transmits light at a different wavelength from that at which it receives light. Therefore, such an optical fiber 925 includes two separate blazed FBGs that couple light into and out of each such optical fiber 925. For example, as illustrated in FIG. 13B, optical fiber 925A includes a first blazed FBG 1315A operating at the transmit wavelength, and a second blazed FBG 1330A operating at the receive wavelength. Optical coupler 1305 includes a corresponding first blazed FBG 1320A operating at the transmit wavelength and a second blazed FBG 1335A operating at the receive wavelength. When a proximal portion 1300 of guidewire 905 is fully inserted into receptacle 1310, blazed FBGs 1320A and 1315A are located substantially adjacent to each other, and blazed FBGs 1335A and 1330A are located substantially adjacent to each other. Similarly, optical fiber 925B and optical coupler 1305 respectively include substantially adjacent transmit FBGs 1315B and 1320B and substantially adjacent receive FBGs 1330B and 1335B.

For additional optoacoustic transducer windows 810 at or near the distal portion 900 or elsewhere along guidewire 905, corresponding additional blazed FBGs may similarly be included on optical coupler 1305 at the appropriate wavelength for transmitting and/or receiving optical energy with respect to such additional optoacoustic transducer windows 810. Moreover, optical coupler 1305 need not be located exactly at the proximal end of the guidewire 905, but may instead be located anywhere near the proximal portion 1300 of the guidewire 905 or even further toward the distal portion 900 of the guidewire 905. Also, alignment of the optical coupler 1305 to guidewire 905 need not be limited to butting guidewire 905 into receptacle 1310 of optical coupler 1305; any other alignment mechanism and/or technique is also included.

7. EXAMPLES OF PROCESS AND CONTROL IMAGING ELECTRONICS

FIG. 14A is a block diagram illustrating generally one example of the imaging guidewire 905 and associated interface components. The block diagram of FIG. 14A includes the imaging guidewire 905, which is coupled by optical coupler 1305 to an optoelectronics module 1400. The optoelectronics module 1400 is coupled to an image processing module 1405 and a user interface 1410 that includes a display providing a viewable still and/or video image of the imaging region near one or more acoustic-to-optical transducers using the acoustically-modulated optical signal received therefrom. In one example, the system 1415 illustrated in the block diagram of FIG. 14A uses an image processing module 1405 and a user interface 1410 that are substantially similar to existing acoustic imaging systems.

11

FIG. 14B is a block diagram illustrating generally another example of the imaging guidewire 905 and associated interface components. In this example, the associated interface components include a tissue (and plaque) characterization module 1420 and an image enhancement module 1425. In this example, an input of tissue characterization module 1420 is coupled to an output from optoelectronics module 1400. An output of tissue characterization module 1420 is coupled to at least one of user interface 1410 or an input of image enhancement module 1425. An output of image enhancement module 1425 is coupled to user interface 1410, such as through image processing module 1405.

In this example, tissue characterization module 1420 processes a signal output from optoelectronics module 1400. In one example, such signal processing assists in distinguishing plaque from nearby vascular tissue. Such plaque can be conceptualized as including, among other things, cholesterol, thrombus, and loose connective tissue that build up within a blood vessel wall. Calcified plaque typically reflects ultrasound better than the nearby vascular tissue, which results in high amplitude echoes. Soft plaques, on the other hand, produce weaker and more texturally homogeneous echoes. These and other differences distinguishing between plaque deposits and nearby vascular tissue are detected using tissue characterization signal processing techniques.

For example, such tissue characterization signal processing may include performing a spectral analysis that examines the energy of the returned ultrasound signal at various frequencies. A plaque deposit will typically have a different spectral signature than nearby vascular tissue without such plaque, allowing discrimination therebetween. Such signal processing may additionally or alternatively include statistical processing (e.g., averaging, filtering, or the like) of the returned ultrasound signal in the time domain. Other signal processing techniques known in the art of tissue characterization may also be applied. In one example, the spatial distribution of the processed returned ultrasound signal is provided to image enhancement module 1425, which provides resulting image enhancement information to image processing module 1405. In this manner, image enhancement module 1425 provides information to user interface 1410 that results in a displaying plaque deposits in a visually different manner (e.g., by assigning plaque deposits a discernable color on the image) than other portions of the image. Other image enhancement techniques known in the art of imaging may also be applied. In a further example, similar techniques are used for discriminating between vulnerable plaque and other plaque, and enhancing the displayed image provide a visual indicator assisting the user in discriminating between vulnerable and other plaque.

8. EXAMPLES OF OPTO-ELECTRONICS MODULE

The opto-electronics module 1400 may include one or more lasers and fiber optic elements. In one example, such as where different transmit and receive wavelengths are used, a first laser is used for providing light to the guidewire 905 for the transmitted ultrasound, and a separate second laser is used for providing light to the guidewire 905 for being modulated by the received ultrasound. In this example, a fiber optic multiplexer couples each channel (associated with a particular one of the guidewire's optical fibers 925) to the transmit and receive lasers and associated optics. This reduces system complexity and costs.

In one example, the sharing of transmit and receive components by multiple guidewire channels is possible at least in

12

part because the acoustic image is acquired over a relatively short distance (e.g., millimeters). The speed of ultrasound in a human or animal body is slow enough to allow for a large number of transmit/receive cycles to be performed during the time period of one image frame. For example, at an image depth (range) of about 2 cm, it will take ultrasonic energy approximately 26 microseconds to travel from the sensor to the range limit, and back. In one such example, therefore, an about 30 microseconds transmit/receive (T/R) cycle is used. In the approximately 30 milliseconds allotted to a single image frame, up to 1,000 T/R cycles can be carried out. In one example, such a large number of T/R cycles per frame allows the system to operate as a phased array even though each sensor is accessed in sequence. Such sequential access of the photoacoustic sensors in the guidewire permits (but does not require) the use of one set of T/R opto-electronics in conjunction with a sequentially operated optical multiplexer.

9. EXAMPLE OF US FOR 3-DIMENSIONAL (3-D) IMAGING

In one example, instead of presenting one 2-D slice of the anatomy, the system is operated to provide a 3-D visual image that permits the viewing of a desired volume of the patient's anatomy or other imaging region of interest. This allows the physician to quickly see the detailed spatial arrangement of structures, such as lesions, with respect to other anatomy. In one example, in which the guidewire 905 includes 30 sequentially-accessed optical fibers having up to 10 photoacoustic transducer windows per optical fiber, $30 \times 10 = 300$ T/R cycles are used to collect the image information from all the transducer windows for one image frame. This is well within the allotted 1,000 such cycles for a range of 2 cm, as discussed above. Thus, such an embodiment allows substantially simultaneous images to be obtained from all 10 transducer windows at of each optical fiber at video rates (e.g., at about 30 frames per second for each transducer window). This allows real-time volumetric data acquisition, which offers a distinct advantage over other imaging techniques. Among other things, such real-time volumetric data acquisition allows real-time 3-D vascular imaging, including visualization of the topology of a blood vessel wall, the extent and precise location of plaque deposits, and, therefore, the ability to identify vulnerable plaque.

10. ALTERNATE EXAMPLE OF ACOUSTIC-TO-OPTICAL RECEIVER

FIG. 15 is a cross-sectional schematic diagram illustrating generally one example of an alternate acoustic-to-optical transducer 1500, which in this example is integrated into an optical fiber 105, including fiber core 105 and fiber cladding 120 and covered by coating 930. In the illustrative example of FIG. 15, transducer 1500 includes a blazed FBG 330 in core 115, a translucent deformable (or empty) region 1505 in cladding 120, and an acoustically-deformable light-reflective surface region 1510 overlaying at least a portion of translucent region 1505. In one example, acoustic-to-optical transducer 1500 is fabricated in a window 810 of an imaging guidewire 805 along with an optical-to-acoustic transducer 325, which generates acoustic energy in a nearby imaging region of interest to be received by acoustic-to-optical transducer 1500.

FIG. 16 is a cross-sectional schematic diagram illustrating generally one example of acoustic-to-optical transducer 1500 in operation. FBG 330 receives light from a proximal end of fiber core 105, and directs the received light outward through

13

translucent region **1505** such that the light impinges upon, and is reflected by, reflective region **1510**. At least some of the reflected light is received at FBG **330** and directed back toward the proximal end of fiber core **105**. As illustrated in FIG. **16**, reflective region **1510** deflects in response to acoustic energy received from the nearby imaging region of interest as a result of insonification by a nearby optical-to-acoustic transducer **325**. The deflection of reflective region **1510** modulates the distance that the light travels between FBG **330** and reflective region **1510**. The resulting change in wavelength or intensity is monitored by interface optoelectronics coupled to a proximal end of optical fiber **105**, such as using the above-described components and techniques.

As illustrated in the example of FIGS. **15** and **16**, acoustic-to-optical transducer **1500** need only include a single FBG (e.g., blazed FBG **330**). Moreover, acoustic-to-optical transducer **1500** need not rely on the Poisson effect in which received acoustic energy “squeezes” in a first direction, thereby modulating an interferometric strain-sensing distance in a second direction that is normal to the first direction. An acoustic-to-optical transducer using the Poisson effect typically suffers from some attenuation in translating the mechanical force from the first direction to the orthogonal second direction. As illustrated in FIG. **16**, however, the acoustic-to-optical transducer **1500** detects a modulating distance that is in the substantially the same direction as the received acoustic energy. Moreover, because region **1510** is reflective, a given deflection results in a modulation of twice the number of wavelengths of light in that deflection distance. This further increases the sensitivity of acoustic-to-optical transducer **1500**.

In one example, region **1505** is filled with a transparent polymer to allow optical energy to pass through. In a further example, region **1505** has a thickness **1515** that is $\frac{1}{4}$ -wave resonant with the received acoustic pressure wave. In such an example, the resonance of the polymer-filled region **1505** serves to increase the motion of the reflective region **1510** over that which would occur if region **1505** were formed of glass. In a further example, the polymer-filled region **1505** includes an acoustic impedance that is close to that of water and, therefore, human or animal tissue.

11. CONCLUSION

Although certain of the above examples have been described with respect to intravascular imaging (e.g., for viewing and/or identifying vulnerable plaque), the present systems, devices, and methods are also applicable to imaging any other body part. For example, for example guidewire or other elongate body as discussed above could be inserted into a biopsy needle, laparoscopic device, or any other lumen or cavity for performing imaging. Moreover, such imaging need not involve insertion of an elongate body into a lumen, for example, an imaging apparatus could alternatively be wrapped around a portion of a region to be imaged.

In another example, this technology can be used to process the Doppler shift in acoustic frequency to image blood flow. The operation would be similar to that described above, however, this would increase the length of the transmitted acoustic signal, and would use known Doppler signal processing in the image processing portion of the control electronics. The transmitted acoustic signal can be lengthened by repeatedly pulsing the transmit optical energy at the same rate as the desired acoustic frequency.

It is to be understood that the above description is intended to be illustrative, and not restrictive. For example, the above-described embodiments may be used in combination with

14

each other. Many other embodiments will be apparent to those of skill in the art upon reviewing the above description. The scope of the invention should, therefore, be determined with reference to the appended claims, along with the full scope of equivalents to which such claims are entitled. In the appended claims, the terms “including” and “in which” are used as the plain-English equivalents of the respective terms “comprising” and “wherein.” Moreover, the terms “first,” “second,” and “third,” etc. are used merely as labels, and are not intended to impose numerical requirements on their objects.

What is claimed is:

1. An imaging apparatus comprising:

an imaging guidewire including:

a first optical fiber including a first blazed fiber Bragg grating (FBG);

an optical coupler defining a receptacle configured to receive the imaging guidewire, the optical coupler including:

a second optical fiber including a second blazed FBG, wherein when the imaging guidewire is positioned within the receptacle, the first blazed FBG is located substantially adjacent the second blazed FBG, and wherein the second blazed FBG is configured to couple light into or out of the first blazed FBG.

2. The imaging apparatus of claim 1,

the imaging guidewire including:

a third optical fiber including a third blazed FBG; and the optical coupler including:

a fourth optical fiber including a fourth blazed FBG, wherein when the imaging guidewire is positioned within the receptacle, the third blazed FBG is located substantially adjacent the fourth blazed FBG, and wherein the fourth blazed FBG is configured to couple light into or out of the third blazed FBG.

3. The imaging apparatus of claim 2,

wherein the third optical fiber includes a fifth blazed FBG, wherein the fourth optical fiber includes a sixth blazed FBG,

wherein when the imaging guidewire is positioned within the receptacle, the fifth blazed FBG is located substantially adjacent the sixth blazed FBG,

wherein the sixth blazed FBG is configured to couple light into or out of the fifth blazed FBG, and

wherein the light coupled into or out of the first blazed FBG has a first wavelength and the light coupled into or out of the fifth blazed FBG has a second wavelength.

4. The imaging apparatus of claim 3,

wherein the first optical fiber includes a seventh blazed FBG,

wherein the second optical fiber includes an eighth blazed FBG,

wherein when the imaging guidewire is positioned within the receptacle, the seventh blazed FBG is located substantially adjacent the eighth blazed FBG, and

wherein the eighth blazed FBG is configured to couple light into or out of the seventh blazed FBG.

5. The imaging apparatus of claim 1, wherein the imaging guidewire has a first end, and

wherein when the first end of the imaging guidewire is positioned within the receptacle, the first blazed FBG is located substantially adjacent the second blazed FBG.

6. The imaging apparatus of claim 1, wherein when the imaging guidewire is positioned within the receptacle includes butting the imaging guidewire into the receptacle.

15

7. An imaging apparatus comprising:
 an imaging guidewire including:
 a first optical fiber including a first blazed fiber Bragg
 grating (FBG);
 a second optical fiber including a second blazed FBG; 5
 an optical coupler defining a receptacle configured to
 receive the imaging guidewire, the optical coupler
 including:
 a third optical fiber including a third blazed FBG, 10
 a fourth optical fiber including a fourth blazed FBG
 wherein when the imaging guidewire is positioned within
 the receptacle:
 the first blazed FBG is located substantially adjacent the
 third blazed FBG, and 15
 the second blazed FBG is located substantially adjacent
 the fourth blazed FBG,
 wherein the third blazed FBG is configured to couple light
 into or out of the first blazed FBG, and the fourth blazed
 FBG is configured to couple light into or out of the 20
 second blazed FBG.

8. An imaging apparatus comprising:
 an imaging guidewire including:
 a first optical fiber including a first blazed fiber Bragg
 grating (FBG) and a second blazed FBG;

16

a second optical fiber including a third blazed FBG and
 a fourth blazed FBG;
 an optical coupler defining a receptacle configured to
 receive the imaging guidewire, the optical coupler
 including:
 a third optical fiber including a fifth blazed FBG and a
 sixth blazed FBG,
 a fourth optical fiber including a seventh blazed FBG
 and an eighth blazed FBG,
 wherein when the imaging guidewire is positioned within
 the receptacle:
 the first and second blazed FBGs are located substan-
 tially adjacent the fifth and sixth blazed FBGs, respec-
 tively, and
 the third and fourth blazed FBGs are located substan-
 tially adjacent the seventh and eighth blazed FBGs,
 respectively,
 wherein one of the fifth and sixth blazed FBGs is config-
 ured to couple light having a first wavelength into or out
 of one of the first and second blazed FBGs, and
 wherein one of the seventh and the eighth blazed FBGs
 is configured to couple light having a second wave-
 length into or out of one of the third and fourth blazed
 FBGs.

* * * * *